SCORE Search Results Details for Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rge.

Score Home Page

Retrieve Application List

SCORE System Overview

SCORE FAQ

Comments / Suggestions

This page gives you Search Results detail for the Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rge.

start

Go Back to previous page

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OM nucleic - nucleic search, using sw model

Run on:

April 25, 2006, 09:55:26; Search time 1771 Seconds

(without alignments)

4012.101 Million cell updates/sec

Title:

US-10-088-319-2_COPY_331_455

Perfect score:

Sequence:

1 caaaaacaaaaacctttac.....tgagtaaggtggccactttg 125

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched:

5883141 seqs, 28421725653 residues

Total number of hits satisfying chosen parameters:

11766282

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

'GenEmbl: * 1: qb ba:* gb in:* gb_env:* 3: 4: gb om:* qb ov:* 6: qb pat:* 7: gb ph:* 8: gb pr:* 9: gb ro:*

10: gb sts:* 11: qb sy:* 12: qb un:* 13: gb vi:* 14: gb htg:* 15: gb_pl:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

			g.			001111111	
Res	ult		Query				
	No.	Score		Length	DB	ID	Description
	- -	124	99.2	652	 8	 D5060 7	D50607 Human CD40
	2	124	99.2	1313	8	HUMCD40L5P	L47983 Homo sapien
	3	124		151978	14	CT005235	CT005235 Pan trogl
	4	124		175706	8	AL135783	AL135783 Human DNA
С	5	115		259175	14	AC106521	AC106521 Rattus no
	6	113.4		205673	14	AC124137	AC124137 Rattus no
	7	112	89.6	2395	6	AR350233	AR350233 Sequence
	8	112	89.6	2395	6	AR629955	AR629955 Sequence
	9	112	89.6	2395	6	AX351057	AX351057 Sequence
	10	112	89.6	6653	8	D31797	D31797 Homo sapien
	11	109.8	87.8	1878	4	AY333790	AY333790 Canis fam
	12	109.8	87.8	1878	6	BD211546	BD211546 Canine an
С	13	109.8	87.8	1878	6	BD211547	BD211547 Canine an
	14	109.8	87.8	1878	6	AR241524	AR241524 Sequence
C	15	109.8	87.8	1878	6	AR241525	AR241525 Sequence
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	20	97	77.6	620	6	E09510	E09510 DNA encodin
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С	30	84.6	67.7	7057	6	AX348840	AX348840 Sequence
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	38 39	32.8 32.8		110000 110000	6	BD426631_07 AR274513 07	Continuation (8 of
					6		Continuation (8 of
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С	43	31.4	23.1	124704	O	WIII 22 I 24	AL159154 Human DNA

ALIGNMENTS.

RESULT 1 D50607

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                                                     linear
                                                              PRI 10-FEB-1999
DEFINITION
           Human CD40 ligand (hCD40L) gene, 5'-flanking region.
ACCESSION
           D50607
           D50607.1 GI:849048
VERSION
KEYWORDS
           CD40 ligand; hCD40L; type-II membrane glycoprotein.
SOURCE
           Homo sapiens (human)
  ORGANISM
           Homo sapiens
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
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REFERENCE
           Seyama, K., Kira, S., Ishidoh, K., Souma, S., Miyakawa, T. and
  AUTHORS
  TITLE
           Genomic structure and PCR-SSCP analysis of the human CD40 ligand
           gene: its application to prenatal screening for X-linked hyper-IgM
           syndrome
  JOURNAL
           Hum. Genet. 97 (2), 180-185 (1996)
   PUBMED
           8566950
REFERENCE
           2
              (bases 1 to 652)
  AUTHORS
           Ishidoh, K.
  TITLE
           Direct Submission
  JOURNAL
           Submitted (19-MAY-1995) Kazumi Ishidoh, Juntendo University School
           of Medicine, Department of Biochemistry; 2-1-1 Hongo, Bunkyo-ku,
           Tokyo 113, Japan (Tel:03-5802-1031, Fax:03-5802-5889)
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                                                     linear
                                                              PRI 06-FEB-2001
DEFINITION Homo sapiens CD40 ligand gene, promoter and partial cds.
           L47983
ACCESSION
VERSION
           L47983.1 GI:1129041
KEYWORDS
SOURCE
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  ORGANISM
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           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
              (bases 1 to 1313)
  AUTHORS
           Schubert, L.A., King, G., Cron, R.Q., Lewis, D.B., Aruffo, A. and
           Hollenbaugh, D.
  TITLE
           The human gp39 promoter. Two distinct nuclear factors of activated
           T cell protein-binding elements contribute independently to
           transcriptional activation
  JOURNAL
           J. Biol. Chem. 270 (50), 29624-29627 (1995)
  PUBMED
           8530342
REFERENCE
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 AUTHORS
           Schubert, L.A., King, G., Cron, R.Q., Lewis, D.B., Aruffo, A. and
           Hollenbaugh, D.
  TITLE
           Direct Submission
  JOURNAL
           Submitted (20-DEC-1995) Dept of Pediatrics, Stanford University,
           300 Pasteur Drive CCSR Bldg Room 2115b, Stanford, CA 94305, USA
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                                                                  Gaps
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Qу
             IIII
Db
        1225 TTTG 1228
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RESULT 3 CT005235 LOCUS CT005235 151978 bp linear DNA HTG 14-JUN-2005 DEFINITION Pan troglodytes chromosome X clone RP43-015P08 map Xq28, *** SEQUENCING IN PROGRESS ***, 5 unordered pieces. ACCESSION CT005235 CT005235.1 GI:67772562 VERSION HTG; HTGS PHASE1. KEYWORDS SOURCE Pan troglodytes (chimpanzee) ORGANISM Pan troglodytes Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Pan. REFERENCE 1 (bases 1 to 151978) AUTHORS Scheer, S., Kuhl, H., Kube, M., Mueller, I., Thiel, J., Borzym, K., Klages, S., Sontag, M., Kosiura, A., Sudbrak, R., Beck, A., Lehrach, H., Yaspo, M.L. and Reinhardt, R. TITLE Direct Submission JOURNAL Submitted (14-JUN-2005) COMMENT ----- Genome Center Center: Max-Planck-Institute for Molecular Genetics Center code: MPIMG ----- Project Information Center clone name: RP43-015P08 ----- Summary Statistics Sequencing vector: pUC19; 100% of reads Chemistry: Dye-terminator Big Dye; 100% of reads Assembly program: Phrap; version 0.990329 Consensus quality: 149183 bases at least Q40 Consensus quality: 149379 bases at least Q30 Consensus quality: 149536 bases at least Q20 Quality coverage: 6.9 This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. ------The RPCI-43 chimpanzee BAC library was prepared from DNA isolated from the blood of a single male chimpanzee using published

The RPCI-43 chimpanzee BAC library was prepared from DNA isolated from the blood of a single male chimpanzee using published protocols (Osoegawa, K. et al. Genomics 51:1-8). The DNA from the chimpanzee ('Clint') was obtained from the Yerkes Primate Center in Atlanta. The library was prepared by Baoli Zhu, Chung Li Shu, Kazutoyo Osoegawa, Evan Eichler & Pieter J de Jong. The library characteristics are described at

http://www.chori.org/bacpac/mchimp43.htm.

The clone may be obtained from Pieter J. de Jong and coworkers (http://www.chori.org/bacpac).

VECTOR: pBACe3.

contig 01 1. .14831 contig 02 14932. .33675 contig 03 33776. .60588 contig 04 60689. .99409 contig 05 99510. .151978.

^{*} NOTE: This is a 'working draft' sequence. It currently

^{*} consists of 5 contigs. The true order of the pieces

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* is not known and their order in this sequence record is
             arbitrary. Gaps between the contigs are represented as
             runs of N, but the exact sizes of the gaps are unknown.
             This record will be updated with the finished sequence
             as soon as it is available and the accession number will
             be preserved.
                         14831: contig of 14831 bp in length
                14832
                         14931: gap of unknown length
                         33675: contig of 18744 bp in length
                14932
                         33775: gap of unknown length
                33676
                33776
                         60588: contig of 26813 bp in length
                60589
                         60688: gap of unknown length
                60689
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Qy
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Qу
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Db
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RESULT 4
AL135783
LOCUS
           AL135783
                                175706 bp
                                             DNA
                                                             PRI 18-MAY-2005
                                                     linear
DEFINITION Human DNA sequence from clone RP3-527F8 on chromosome Xq25-27.1
           Contains the TNFSF5 gene for tumor necrosis factor (ligand)
           superfamily (member 5 , hyper-IgM syndrome), a novel gene, the 3'
           end of the ARHGEF6 gene for Rac/Cdc42 guanine nucleotide exchange
           factor (GEF) 6 and a CpG island, complete sequence.
ACCESSION
           AL135783
           AL135783.6 GI:6983480
VERSION
KEYWORDS
           HTG; ARHGEF6; TNFSF5.
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SOURCE Homo sapiens (human) ORGANISM Homo sapiens Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 175706) REFERENCE AUTHORS Bird, C. TITLE Direct Submission JOURNAL Submitted (13-MAY-2005) Wellcome Trust Sanger Institute, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: vega@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk On Feb 16, 2000 this sequence version replaced gi:6983044. COMMENT The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome X, constructed by the Sanger Centre Chromosome X Mapping Group. Further information can be found at http://www.sanger.ac.uk/HGP/ChrX ----- Genome Center Center: Wellcome Trust Sanger Institute Center code: SC Web site: http://www.sanger.ac.uk Contact: vega@sanger.ac.uk This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one subclone; and the assembly was confirmed by restriction digest, except on the rare occasion of the clone being a YAC. RP3-527F8 is from the library RPCI-3 constructed by the group of Pieter de Jong. For further details see http://www.chori.org/bacpac/home.htm VECTOR: pCYPAC2. **FEATURES** Location/Qualifiers source 1. .175706 /organism="Homo sapiens" /mol type="genomic DNA" /db xref="taxon:9606" /map = "q25 - 27.1"/clone="RP3-527F8" /clone_lib="RPCI-3" 39996. .42685 gene /locus tag="RP3-527F8.2-001" mRNA join(39996. .40060,41030. .42685) /locus tag="RP3-527F8.2-001" /product="novel protein" /note="match: cDNAs: BC042469.1" CDS 41116. .41343 /locus tag="RP3-527F8.2-001" /standard_name="OTTHUMP00000024129" /codon start=1 /product="novel protein" /protein_id="CAI42900.1" /db_xref="GI:57208602" /db xref="UniProt/TrEMBL:Q5JVP7" translation="MSSSLWVLPHASSCTWNVFFPDPENSFRASSSRKSSLSLSPLHP/

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LOCUS
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DEFINITION
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            ***, 3 unordered pieces.
            AC106521
ACCESSION
VERSION
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KEYWORDS
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SOURCE
           Rattus norvegicus (Norway rat)
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REFERENCE
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 AUTHORS
           Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J.,
            Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D.,
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Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokelemeh, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkoch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A. Direct Submission Unpublished 2 (bases 1 to 259175) Worley, K.C. Direct Submission Submitted (12-JAN-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA 3 (bases 1 to 259175) Rat Genome Sequencing Consortium. Direct Submission Submitted (08-OCT-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA On Sep 14, 2002 this sequence version replaced gi:21731842. The sequence in this assembly is a combination of BAC based reads and whole genome shotgun sequencing reads assembled using Atlas (http://www.hgsc.bcm.tmc.edu/projects/rat/). Each contig described in the feature table below represents a scaffold in the Atlas assembly (a 'contig-scaffold'). Within each contig-scaffold, individual sequence contigs are ordered and oriented, and separated by sized gaps filled with Ns to the estimated size. The sequence may extend beyond the ends of the clone and there may be sequence contigs within a contig-scaffold that consist entirely of whole genome shotgun sequence reads. Both end sequences and whole genome shotgun sequence only contigs will be indicated in the feature table. ----- Genome Center

Center: Baylor College of Medicine

Center code: BCM

TITLE

REFERENCE AUTHORS

TITLE

REFERENCE

TITLE

COMMENT

AUTHORS

JOURNAL

JOURNAL

JOURNAL

Web site: http://www.hgsc.bcm.tmc.edu/

Contact: hgsc-help@bcm.tmc.edu ----- Project Information

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Center project name: GLCE
               Center clone name: CH230-29H24
              ----- Summary Statistics
               Assembly program: Phrap; version 0.990329
               Consensus quality: 236111 bases at least Q40
               Consensus quality: 238370 bases at least Q30
               Consensus quality: 239565 bases at least Q20
               Estimated insert size: 259108; sum-of-contigs estimation
               Quality coverage: 4x in Q20 bases; sum-of-contigs estimation
             NOTE: Estimated insert size may differ from sequence length
               (see http://www.hgsc.bcm.tmc.edu/docs/Genbank draft data.html)
            NOTE: This sequence may represent more than one clone.
            * NOTE: This is a 'working draft' sequence. It currently
            * consists of 3 contigs. The true order of the pieces
           * is not known and their order in this sequence record is
             arbitrary. Gaps between the contigs are represented as
             runs of N, but the exact sizes of the gaps are unknown.
             This record will be updated with the finished sequence
             as soon as it is available and the accession number will
             be preserved.
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Qу
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RESULT 6
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LOCUS
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                                            DNA
                                                   ·linear
                                                             HTG 20-NOV-2002
DEFINITION Rattus norvegicus clone CH230-285J20, WORKING DRAFT SEQUENCE.
ACCESSION AC124137
VERSION
           AC124137.4 GI:25139769
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HTG; HTGS PHASE2; HTGS DRAFT; HTGS FULLTOP. SOURCE Rattus norvegicus (Norway rat) ORGANISM Rattus norvegicus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi; Muroidea; Muridae; Murinae; Rattus. REFERENCE (bases 1 to 205673) **AUTHORS** Muzny, D. Marie., Metzker, M. Lee., Abramzon, S., Adams, C., Alder, J., Allen, C., Allen, H., Alsbrooks, S., Amin, A., Anguiano, D., Anyalebechi, V., Aoyagi, A., Ayodeji, M., Baca, E., Baden, H., Baldwin, D., Bandaranaike, D., Barber, M., Barnstead, M., Benahmed, F., Biswalo, K., Blair, J., Blankenburg, K., Blyth, P., Brown, M., Bryant, N., Buhay, C., Burch, P., Burrell, K., Calderon, E., Cardenas, V., Carter, K., Cavazos, I., Ceasar, H., Center, A., Chacko, J., Chavez, D., Chen, G., Chen, R., Chen, Y., Chen, Z., Chu, J., Cleveland, C., Cockrell, R., Cox, C., Coyle, M., Cree, A., D'Souza, L., Davila, M.L., Davis, C., Davy-Carroll, L., De Anda, C., Dederich, D., Delgado, O., Denson, S., Deramo, C., Ding, Y., Dinh, H., Divya, K., Draper, H., Dugan-Rocha, S., Dunn, A., Durbin, K., Duval, B., Eaves, K., Egan, A., Escotto, M., Eugene, C., Evans, C.A., Falls, T., Fan, G., Fernandez, S., Finley, M., Flagg, N., Forbes, L., Foster, M., Foster, P., Fraser, C.M., Gabisi, A., Ganta, R., Garcia, A., Garner, T., Garza, M., Gebregeorgis, E., Geer, K., Gill, R., Grady, M., Guerra, W., Guevara, W., Gunaratne, P., Haaland, W., Hamil, C., Hamilton, C., Hamilton, K., Harvey, Y., Havlak, P., Hawes, A., Henderson, N., Hernandez, J., Hernandez, R., Hines, S., Hladun, S.L., Hodgson, A., Hoques, M., Hollins, B., Howells, S., Hulyk, S., Hume, J., Idlebird, D., Jackson, A., Jackson, L., Jacob, L., Jiang, H., Johnson, B., Johnson, R., Jolivet, A., Karpathy, S., Kelly, S., Kelly, S., Khan, Z., King, L., Kovar, C., Kowis, C., Kraft, C.L., Lebow, H., Levan, J., Lewis, L., Li, Z., Liu, J., Liu, J., Liu, W., Liu, Y., London, P., Longacre, S., Lopez, J., Lorensuhewa, L., Loulseged, H., Lozado, R.J., Lu, X., Ma, J., Maheshwari, M., Mahindartne, M., Mahmoud, M., Malloy, K., Mangum, A., Mangum, B., Mapua, P., Martin, K., Martin, R., Martinez, E., Mawhiney, S., McLeod, M.P., McNeill, T.Z., Meenen, E., Milosavljevic, A., Miner, G., Minja, E., Montemayor, J., Moore, S., Morgan, M., Morris, K., Morris, S., Munidasa, M., Murphy, M., Nair, L., Nankervis, C., Neal, D., Newton, N., Nguyen, N., Norris, S., Nwaokelemeh, O., Okwuonu, G., Olarnpunsagoon, A., Pal, S., Parks, K., Pasternak, S., Paul, H., Perez, A., Perez, L., Pfannkoch, C., Plopper, F., Poindexter, A., Popovic, D., Primus, E., Pu, L.-L., Puazo, M., Quiroz, J., Rachlin, E., Reeves, K., Regier, M.A., Reigh, R., Reilly, B., Reilly, M., Ren, Y., Reuter, M., Richards, S., Riggs, F., Rives, C., Rodkey, T., Rojas, A., Rose, M., Rose, R., Ruiz, S.J., Sanders, W., Savery, G., Scherer, S., Scott, G., Shatsman, S., Shen, H., Shetty, J., Shvartsbeyn, A., Sisson, I., Sitter, C.D., Smajs, D., Sneed, A., Sodergren, E., Song, X.-Z., Sorelle, R., Sosa, J., Steimle, M., Strong, R., Sutton, A., Svatek, A., Tabor, P., Taylor, C., Taylor, T., Thomas, N., Thomas, S., Tingey, A., Trejos, Z., Usmani, K., Valas, R., Vera, V., Villasana, D., Waldron, L., Walker, B., Wang, J., Wang, Q., Wang, S., Warren, J., Warren, R., Wei, X., White, F., Williams, G., Willson, R., Wleczyk, R., Wooden, H., Worley, K., Wright, D., Wright, R., Wu, J., Yakub, S., Yen, J., Yoon, L., Yoon, V., Yu, F., Zhang, J., Zhou, J., Zhou, X., Zhao, S., Dunn, D., von Niederhausern, A., Weiss, R., Smith, D.R., Holt, R.A., Smith, H.O., Weinstock, G. and Gibbs, R.A. TITLE Direct Submission JOURNAL Unpublished REFERENCE (bases 1 to 205673) 2 AUTHORS Worley, K.C. TITLE Direct Submission

KEYWORDS .

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Submitted (10-JUN-2002) Human Genome Sequencing Center, Department
  JOURNAL
            of Molecular and Human Genetics, Baylor College of Medicine, One
            Baylor Plaza, Houston, TX 77030, USA
REFERENCE
            3 (bases 1 to 205673)
  AUTHORS
            Rat Genome Sequencing Consortium.
  TITLE
            Direct Submission
            Submitted (20-NOV-2002) Human Genome Sequencing Center, Department
  JOURNAL
            of Molecular and Human Genetics, Baylor College of Medicine, One
            Baylor Plaza, Houston, TX 77030, USA
            On Nov 20, 2002 this sequence version replaced gi:23269442.
COMMENT
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            assembly (a 'contig-scaffold'). Within each contig-scaffold,
            individual sequence contigs are ordered and oriented, and separated
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            may extend beyond the ends of the clone and there may be sequence
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                Center: Baylor College of Medicine
                Center code: BCM
                Web site: http://www.hgsc.bcm.tmc.edu/
                Contact: hqsc-help@bcm.tmc.edu
             ----- Project Information
                Center project name: KABZ
               Center clone name: CH230-285J20
            ----- Summary Statistics
               Assembly program: Phrap; version 0.990329
                Consensus quality: 192345 bases at least Q40
                Consensus quality: 195056 bases at least Q30
                Consensus quality: 196514 bases at least Q20
                Estimated insert size: 198117; sum-of-contigs estimation
                Quality coverage: 7x in Q20 bases; sum-of-contigs estimation
            * NOTE: Estimated insert size may differ from sequence length
                (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft data.html).
            * NOTE: This is a 'working draft' sequence. It currently
             consists of 1 contigs. Gaps between the contigs
            * are represented as runs of N. The order of the pieces
            * is believed to be correct as given, however the sizes
            * of the gaps between them are based on estimates that have
            * provided by the submittor.
            * This sequence will be replaced
            * by the finished sequence as soon as it is available and
            * the accession number will be preserved.
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VERSION
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          Unclassified.
REFERENCE
          1 (bases 1 to 2395)
 AUTHORS
          Bennett, C.F., Baker, B.F., Wyatt, J. and Davis, S.E.
  TITLE
          Antisense modulation of CD40 ligand expression
  JOURNAL
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Qу
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                                                          PAT 14-FEB-2005
DEFINITION Sequence 9 from patent US 6838556.
ACCESSION
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VERSION
          AR629955.1 GI:59762089
KEYWORDS
SOURCE
          Unknown.
 ORGANISM Unknown.
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Unclassified.
REFERENCE
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           Kim, J.P., Starr, D.B., Tam, A.W., Laurance, M.E., Michelotti, E.F.,
  AUTHORS
           Velligan, M.D., Latour, D.R., Thomas, R.L., Kongpachith, A.,
           Sheppard, L.T., Kim, M.Y. and Bruice, T.W.
           Promoters for regulated gene expression
  TITLE
  JOURNAL
           Patent: US 6838556-A 9 04-JAN-2005;
           Genelabs Technologies, Inc.; Redwood City, CA
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 Matches 123; Conservative
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DEFINITION Sequence 9 from Patent W00194600.
ACCESSION
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VERSION
           AX351057.1 GI:18616411
KEYWORDS
SOURCE
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 ORGANISM
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           Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
           Hominidae; Homo.
REFERENCE
 AUTHORS
           Kim, J.P., Starr, D.B., Tam, A.W., Laurance, M.E., Michelotti, E.F.,
           Velligan, M.D., Latour, D.R., Thomas, R.L., Kongpachith, A.,
           Sheppard, L.T., Lim, M.Y. and Bruice, T.W.
 TITLE
           Promoters for regulated gene expression
 JOURNAL
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REFERENCE
  AUTHORS
           Hollenbaugh, D., Grosmaire, L.S., Kullas, C.D., Chalupny, N.J.,
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            and Aruffo, A.
  TITLE
            The human T cell antigen gp39, a member of the TNF gene family, is
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  JOURNAL
           EMBO J. 11 (12), 4313-4321 (1992)
  PUBMED
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REFERENCE
 AUTHORS
           Spriggs, M.K., Armitage, R.J., Strockbine, L., Clifford, K.N.,
           Macduff, B.M., Sato, T.A., Maliszewski, C.R. and Fanslow, W.C.
           Recombinant human CD40 ligand stimulates B cell proliferation and
  TITLE
           immunoglobulin E secretion
  JOURNAL
           J. Exp. Med. 176 (6), 1543-1550 (1992)
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           Villa, A., Noterengelo, L., Disanto, J., Nacchi, P., Strina, D.,
           Frattini, A., Lucchini, F., Patrosso, C., Giliani, S., Mantuano, E.,
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  TITLE
           Organization of the human CD40L gene: implications for molecular
           defects in X chromosome-linked hyper-IgM syndrome and prenatal
           diagnosis
  JOURNAL
           Proc. Natl. Acad. Sci. U.S.A. 91 (6), 2110-2114 (1994)
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REFERENCE
           Shimadzu, M., Nunoi, H., Terasaki, H., Ninomiya, R., Iwata, M.,
 AUTHORS
           Kanegasaka, S. and Matsuda, I.
 TITLE
           Structural organization of the gene for CD40 ligand: molecular
           analysis for diagnosis of X-linked hyper-IqM syndrome
 JOURNAL
           Biochim. Biophys. Acta 1260 (1), 67-72 (1995)
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           Shimadzu, M., Terasaki, H., Ninomiya, R., Shimizu, S., Nunoi, H. and
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TITLE
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  AUTHORS
            Shimadzu, M.
  TITLE
            Direct Submission
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  JOURNAL
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REFERENCE
           1 (bases 1 to 1878)
 AUTHORS
           Yang, S. and Sim, G.-K.
 TITLE
           Canine CD40 and CD40 Ligand cDNA Sequences
 JOURNAL Unpublished
REFERENCE
           2 (bases 1 to 1878)
 AUTHORS
           Yang, S. and Sim, G.-K.
 TITLE
           Direct Submission
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 AUTHORS
          Sim, G., Yang, S., Dreitz, M.J. and Wonderling, R.S.
 TITLE
          Canine and feline immunoregulatory proteins, nucleic acid molecules
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 AUTHORS
          Sim, G., Yang, S., Dreitz, M.J. and Wonderling, R.S.
 TITLE
          Canine and feline immunoregulatory proteins, nucleic acid molecules
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          Patent: JP 2002516104-A 53 04-JUN-2002;
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REFERENCE
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 AUTHORS
          Sim, G.-K., Yang, S., Dreitz, M.J. and Wonderling, R.S.
 TITLE
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 JOURNAL
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 $\mathbf{H}\mathbf{H}\mathbf{I}$

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Job time : 1775 secs

Qу

Db

SCORE 1.3 BuildDate: 12/06/2005

SCORE Search Results Details for Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rng.

Score Home Page

Retrieve Application

SCORE System Overview

SCORE FAQ

Comments / <u>Suggestions</u>

This page gives you Search Results detail for the Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rng.

start

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OM nucleic - nucleic search, using sw model

Run on:

April 25, 2006, 09:28:13; Search time 285 Seconds

(without alignments)

2923.111 Million cell updates/sec

Title:

US-10-088-319-2 COPY 331 455

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Minimum DB seg length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a

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	7	97	77.6	620	2	AAT05790	Aat05790 Human CD4
	8	93.6	74.9	7057	4	AAS46601	Aas46601 Tumour su
	9	93.6	74.9	7057	6	ABL33848	Abl33848 Human imm
	10	93.6	74.9	7057	6	ABL34594	Abl34594 Human met
	11	93.6	74.9	7057	6	ABL70407	Ab170407 Chemicall
	12	93.6	74.9	7057	6	AAS61354	Aas61354 Human gen
	13	93.6	74.9	7057	7	ADS99855	Ads99855 Bisulphit
С	14	84.6	67.7	7057	4	AAS46602	Aas46602 Tumour su
С	15	84.6	67.7	7057	6	ABL33849	Abl33849 Human imm
С	16	84.6	67.7	7057	6	ABL34595	Abl34595 Human met
С	17	84.6	67.7	7057	6	ABL70408	Ab170408 Chemicall
С	18	84.6	67.7	7057	6	AAS,61355	Aas61355 Human gen
C.	19	84.6	67.7	7057	7	ADS99856	Ads99856 Complemen
	20	37	29.6	37	. 6	ABK29866	Abk29866 CD40L -77
	21	32.8		110000	2	AAT42063_07	Continuation (8 of
	22	31.4		290040	14	ADV16961	Adv16961 Human pro
	23	31.4		290040	14	· ADU92049	Adu92049 Human PAM
	24	31	24.8	438	4	AAI87513	Aai87513 Human pol
С	25	30.8	24.6	1470	3	AAC46273	Aac46273 Arabidops
С	26	30.8	24.6	1493	13	ADT15110 ·	Adt15110 Plant cDN
С	27	30.4	24.3	454	5	ABV09654	Abv09654 Human pro
С	28	30.2	24.2	1833	10	ADD48377 .	Add48377 Human gen
С	29	30	24.0	7005	13	ADS47818	Ads47818 Bacterial
	30	29.8	23.8	2004	7	ADZ74714	Adz74714 Arabidops
С	31	29.8	23.8	52679	11	ACN44216	Acn44216 Mouse gen
	32	29.6		130312	14	AEB39168	Aeb39168 L. pneumo
С	33	29.6		143354	14	AEB42740	Aeb42740 L. pneumo
	34	29.4	23.5	1716	4	AAF61006	Aaf61006 P. putida
	35	29.4	23.5	5771	10	ADD45213	Add45213 Rat gene
	36	29.2	23.4	474	4	ABA59013	Aba59013 Human foe
	37	29.2	23.4	474	4	AAI38738	Aai38738 Probe #74
	38	29.2	23.4	474	4	AAK32932	Aak32932 Human bon
	39	29.2	23.4	474	4	AAK07183	Aak07183 Human bra
	40	29.2	23.4		4	ABS32660	Abs32660 Human liv
	41	29.2	23.4	474	6	ABS07737	Abs07737 Human gen
С	42	29.2	23.4	1474	3	AAC33070	Aac33070 Arabidops
	43	29.2	23.4	10929	13	ADT66569	Adt66569 Rat LutzP
	44	29.2	23.4	10929	14		Adx26167 Novel cel
С	45	28.8	23.0	5332	4	ABL29632	Abl29632 Drosophil

ALIGNMENTS

RESULT 1 AAF74905

AAF74905 standard; DNA; 455 BP.

```
XX
     AAF74905;
AC
XX
DT
     23-MAY-2001 (first entry)
XX
     Human altered CD40L promoter sequence (A331C) SEQ ID NO:2.
DΕ
XX
KW
     Human; CD40L; promoter; CD40 ligand promoter; rheumatoid arthritis;
KW
     diagnosis; antiarthritic; antirheumatic; immunosuppressive;
KW
     antiinflammatory; inflammatory disease; autoimmune disease; ds.
XX
os
     Homo sapiens.
os
     Synthetic.
XX
PN
     WO200119844-A1.
XX
PD
     22-MAR-2001.
XX
     13-SEP-2000; 2000WO-US024966.
PF
XX
PR
     13-SEP-1999;
                   99US-0153625P.
XX
PA
     (NYRE-) NEW YORK SOC RELIEF RUPTURED & CRIPPLED.
XX
PΙ
     Crow MK,
             Li Y;
XX
DR
     WPI; 2001-244776/25.
XX
PT
     New altered CD40L promoter for use in the study, diagnosis and treatment
PT
     of a variety of inflammatory disorders and autoimmune diseases, such as
PT
     rheumatoid arthritis.
XX
PS
     Claim 1; Fig 2; 90pp; English.
XX
CC
     The present invention describes an isolated, purified nucleic acid, which
CC .
     is an altered CD40 ligand (CD40L) promoter (I) for CD40 ligand, having
CC
     residues 331-455 of the sequence comprising 455 nucleotides given in
     AAF74905 where A in the wild type sequence at position 331 (corresponding
CC
CC
    to position -125) is replaced with C. (I) has antiarthritic,
CC
     antirheumatic, immunosuppressive and antiinflammatory activities, and can
     be used in gene therapy. (I) is useful in the study, diagnosis and
CC
     treatment of inflammatory and autoimmune diseases, as well as diseases in
CC
     which elevated expression of CD40L is a factor, e.g., rheumatoid
CC
     arthritis. The present sequence represents the specifically claimed
CC
     altered human CD40L promoter sequence, from the present invention
XX
    Sequence 455 BP; 156 A; 72 C; 95 G; 132 T; 0 U; 0 Other;
  Query Match
                         100.0%; Score 125; DB 4; Length 455;
  Best Local Similarity
                         100.0%;
                                 Pred. No. 1.9e-34;
 Matches 125; Conservative
                               0; Mismatches
                                                 0:
                                                     Indels
                                                              0;
                                                                  Gaps
           1 CAAAAACAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTT 60
Qy
             Db
         331 CAAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTT 390
          61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 120
Qy
             Db
         391 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 450
         121 CTTTG 125
Qу
```

```
RESULT 2
AAF74904
     AAF74904 standard; DNA; 455 BP.
XX
AC
     AAF74904;
XX
DT
     23-MAY-2001
                  (first entry)
XX
DE
     Human wild type CD40L promoter sequence SEQ ID NO:1.
XX
KW
     Human; CD40L; promoter; CD40 ligand promoter; rheumatoid arthritis;
KW
     diagnosis; antiarthritic; antirheumatic; immunosuppressive;
KW
     antiinflammatory; inflammatory disease; autoimmune disease; ds.
XX
OS
     Homo sapiens.
XX
     WO200119844-A1.
PN
XX
PD
     22-MAR-2001.
XX
     13-SEP-2000; 2000WO-US024966.
PF
XX
                    99US-0153625P.
PR
     13-SEP-1999;
XX
PA
     (NYRE-) NEW YORK SOC RELIEF RUPTURED & CRIPPLED.
XX
PΙ
     Crow MK,
             Li Y;
XX
     WPI; 2001-244776/25.
DR
XX
PT
     New altered CD40L promoter for use in the study, diagnosis and treatment
     of a variety of inflammatory disorders and autoimmune diseases, such as
PT
     rheumatoid arthritis.
XX
PS
     Example 1; Fig 2; 90pp; English.
XX
CC
     The present invention describes an isolated, purified nucleic acid, which
·CC
     is an altered CD40 ligand (CD40L) promoter (I) for CD40 ligand, having
CC
     residues 331-455 of the sequence comprising 455 nucleotides given in
CC
     AAF74905 where A in the wild type sequence at position 331 (corresponding
CC
     to position -125) is replaced with C. (I) has antiarthritic,
CC
     antirheumatic, immunosuppressive and antiinflammatory activities, and can
CC
     be used in gene therapy. (I) is useful in the study, diagnosis and
CC
     treatment of inflammatory and autoimmune diseases, as well as diseases in
CC
     which elevated expression of CD40L is a factor, e.g., rheumatoid
CC
     arthritis. The present sequence represents the wild type human CD40L
CC
     promoter sequence, which is used in an example from the present invention
XX
SQ
     Sequence 455 BP; 157 A; 71 C; 95 G; 132 T; 0 U; 0 Other;
                         99.2%; Score 124; DB 4; Length 455;
  Query Match
                         100.0%;
  Best Local Similarity
                                 Pred. No. 4.3e-34;
  Matches 124; Conservative
                                0; Mismatches
                                                      Indels
                                                                            0;
                                                  0;
                                                                0;
                                                                    Gaps
            2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
Qу
              Db
          332 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 391
```

```
62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qy
              Db
         392 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 451
         122 TTTG 125
Qу
              IIII
Db
         452 TTTG 455
RESULT 3
ABK29860
    ABK29860 standard; DNA; 2395 BP.
TD
XX
AC
    ABK29860;
XX
DΤ
     23-APR-2002 (first entry)
XX
DE
    Wild type CD40Ln D1 promoter, nucleotides -1860 to +49.
XX
KW
    Cyclin D1 promoter; CD40L promoter; hepatitis B virus promoter;
KW
     HBV promoter; vancomycin-resistant enterococci promoter; VRE promoter;
    vanH promoter; androgen receptor promoter; AR promoter;
KW
KW
     human epidermal growth factor receptor 2 promoter; her2 promoter;
KW
    beta lactamase promoter; Bla promoter; transgene; cancer; breast cancer;
KW
     colon cancer; immunological disorder; prostate cancer; cytostatic;
KW
     autoimmune disease; HBV pre-S promoter; HBV-X promoter;
KW
    Enterococcus infection; immunosuppressive; antibacterial; antiviral;
KW
     gene expression modulator; multiple sclerosis; MS;
KW
     chronic hepatic insufficiency; cirrhosis; hepatocellular carcinoma;
KW
     systematic lupus erythematosus; SLE; graft-vs-host disease; GVHD;
KW
     familial adenomatous polyposis; rheumatoid arthritis; PCR; primer;
KW
     transgenic; ds.
XX
os
    Homo sapiens.
XX
    W0200194600-A2.
PN
XX
PD
    13-DEC-2001.
XX
PF
     06-JUN-2001; 2001WO-US018343.
XX
     06-JUN-2000; 2000US-0209549P.
PR
XX
PA
     (GENE-) GENELABS TECHNOLOGIES INC.
XX
PΙ
    Kim JP, Starr DB, Tam AW, Laurance ME, Michelotti EF;
ΡI
    Velligan MD, Latour DR, Thomas RL, Kongpachith A, Sheppard LT;
PΙ
    Lim MY, Bruice TW;
XX
DR
    WPI; 2002-130595/17.
XX
PT
    New nucleic acid regulatory sequences, which are able to regulate
PT
     expression of a gene operably linked to a promoter, useful for regulating
PT
     the expression of transgenes and for treating e.g., cancer and
PT
    immunological diseases.
XX
PS
    Example 2; Fig 5A-C; 95pp; English.
XX
CC
    The invention describes an isolated nucleic acid regulatory sequence for
     a cyclin D1 promoter, a CD40L promoter, vancomycin-resistant enterococci
```

```
CC
     (VRE) promoter, an HBV promoter, androgen receptor (AR) promoter, Human
     epidermal growth factor receptor 2 (HER2) promoter, or a beta lactamase
CC
CC
     (Bla) promoter. Transcription regulatory sequences may be used to
CC
     regulate expression of the endogenous, autologous or heterologous genes
     operably linked to the promoter, and may be incorporated into
CC
CC
     heterologous nucleic acid constructs for use in regulated expression of
CC
     transgenes. Regulated expression of cyclin D1 can be used in cancer
CC
     therapies, such as breast, colon or pancreatic cancers and familial
     adenomatous polyposis. Regulation of the activity of CD40L gene promoter
CC
CC
     may be used in the treatment of immunological disorders, such as
CC
     autoimmune diseases e.g. multiple sclerosis (MS), systematic lupus
CC
     erythematosus (SLE), graft-vs-host disease (GVHD) and rheumatoid
CC
     arthritis. Regulated expression of genes under the control of the HBV
CC
     (hepatitis B)-specific core, pre-S and X promoters can be used in the
CC
     therapy of HBV disease, chronic hepatic insufficiency, cirrhosis,
CC
     hepatocellular carcinoma, and in the regulated expression of liver cell-
     specific genes. Regulated expression of the vanH gene promoter can be
CC
CC
     used in treatment of Enterococcus infection, while regulated expression
CC
     of the androgen receptor gene can be used in the treatment of prostate
CC
     cancer. This sequence represents the human CD40L promoter, nucleotides -
     1860 to +49, the regulatory regions of which are described in the method
CC
CC
     of the invention
XX.
so
     Sequence 2395 BP; 708 A; 453 C; 502 G; 732 T; 0 U; 0 Other;
  Query Match
                         89.6%;
                                 Score 112; DB 6; Length 2395;
  Best Local Similarity
                         99.2%;
                                 Pred. No. 1.6e-29;
 Matches 123; Conservative
                                0; Mismatches
                                                 0;
                                                     Indels
                                                                   Gaps
                                                                           1:
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
Qу
             Db
        1750 AAAAACAAAAACCTTTACGTAACG-TTTTGCTGGGAGAGAGACTACGAAGCACATTTT 1808
Qу
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
             1809 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 1868
Db
         122 TTTG 125
Qу
             Db
        1869 TTTG 1872
RESULT 4
ACC44010
     ACC44010 standard; DNA; 2395 BP.
XX
AC
    ACC44010;
XX
DT
     30-MAY-2003 (first entry)
XX
DE
    Human CD40 ligand gene fragment #1.
XX
KW
     ds; cytostatic; antiinflammatory; immunomodulator; antisense; gene;
KW
     gene therapy; human; CD40 ligand; phosphorothioate; 2'MOE wings; cancer;
     autoimmune disorder; inflammatory disorder; apoptosis.
KW
XX
os
     Homo sapiens.
XX
FH
                    Location/Qualifiers
    Key
FT
                    1939. .2094
     CDS
FT
                    /*tag= a
```

```
FT
                    /product='"CD40 ligand fragment"
FT
                    /note= "no stop codon given"
XX
PN
    WO2003008433-A1.
XX
PD
    30-JAN-2003.
XX
    15-JUL-2002; 2002WO-US022635.
PF
XX
    18-JUL-2001; 2001US-00909595.
PR
XX
     (ISIS-) ISIS PHARM INC.
PΑ
XX
PΙ
    Bennett CF, Baker BF, Wyatt JR, Davis SE;
XX
DR
    WPI; 2003-239305/23.
DR
    P-PSDB; ABP98669.
XX
PT
    New antisense oligonucleotides targeted to nucleic acids encoding a CD40
    ligand, useful in diagnostic and research applications, or for treating
PT
    diseases associated with expression of CD40 ligand, e.g. cancer or
PT
    autoimmune disorder.
XX
PS
    Example 13; Page 87-89; 108pp; English.
XX
CC
    The invention relates to novel antisense oligonucleotide targeted to the
CC
    human CD40 ligand gene. The oligonucleotides contain either
    phosphorothicate internucleotide bonds replacing the usual phosphodiester
CC
CC
    internucleotide bonds or have a peptide amide backbone replacing the
CC
    sugar phosphate backbone. The nucleotides flanking the central 10
    nucleotides have 2'-methoxyethyl nucleotides (2'MOE wings) and the
CC
CC
    cytidine nucleotides are all 5-methylcytidines. The antisense compounds
    are useful for modulating the expression of CD40 ligand and for treating
CC
CC
    diseases or conditions associated with expression of CD40 liqand, e.q.
CC
    cancer, autoimmune disorder, inflammatory disorder, or a disease or
CC
    condition arising from aberrant apoptosis. The antisense compounds are
    also useful for diagnostics, therapeutics, prophylaxis, e.g. to prevent
CC
CC
    or delay infection, inflammation or tumor formation, as research reagents
CC
    and kits, and in distinguishing between functions of various members of a
CC
    biological pathway. This sequence represents a fragment of the genomic
CC
    sequence encoding the human CD40 ligand. The sequence was used to design
CC
    the novel antisense oligonucleotides of the invention
XX
so
    Sequence 2395 BP; 708 A; 453 C; 502 G; 732 T; 0 U; 0 Other;
                         89.6%;
 Query Match
                                Score 112; DB 8; Length 2395;
 Best Local Similarity
                        99.2%; Pred. No. 1.6e-29;
 Matches 123; Conservative
                               0; Mismatches
                                                    Indels
Qy
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61
             1750 AAAAACAAAAAACCTTTACGTAACG-TTTTGCTGGGAGAGACTACGAAGCACATTTT 1808
Db
Qy
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
             Db
        1809 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 1868
         122 TTTG 125
Qу
             1111
Db
        1869 TTTG 1872
```

```
RESULT 5
AAZ55534
     AAZ55534 standard; cDNA; 1878 BP.
ID
XX
AC.
     AAZ55534;
XX
     14-MAR-2000 (first entry)
DΤ
XX
     Canine CD154 cDNA.
DE
XX
     CD154; CD40 ligand; antibody; canine; feline; inhibitor; immune response;
KW
KW
     immunoregulation; tumour; cancer; autoimmune disease; vaccine; ss.
XX
OS
     Canis familiaris.
XX
FH
                     Location/Qualifiers
     Key
FT
     CDS
                     284. .1066
FT
                     /*tag= a
FT
                     /product= "Canine CD154"
XX
PN
     W09961618-A2.
XX
     02-DEC-1999.
PD
XX
PF
     28-MAY-1999;
                    99WO-US011942.
XX
                    98US-0087306P.
PR
     29-MAY-1998;
XX
     (HESK-) HESKA CORP.
PA
XX
     Sim G, Yang S, Dreitz MJ, Wonderling RS;
PΙ
XX
DR
     WPI; 2000-072623/06.
DR
     P-PSDB; AAY58215.
XX
PT
     Nucleic acids encoding immunoregulatory proteins from cats or dogs,
PΨ
     useful for treating or preventing e.g. tumors or autoimmune disease.
XX
PS
     Claim 1f; Page 205-207; 264pp; English.
XX
     Sequences AAZ55533-Z55539 represent full-length or partial cDNAs encoding
CC
CC
     canine CD154 (CD40 ligand), while sequences AAZ55540-Z55545 represent
     feline CD154 cDNA sequences. The invention relates to canine interleukin-
CC
CC
     4 (IL-4), canine or feline Flt-3 ligand, canine or feline CD40, canine or
CC
     feline CD154, canine IL-5, canine IL-13, feline interferon-alpha (IFN-
CC
     alpha) and feline granulocyte macrophage colony-stimulating factor
CC
     (GMCSF), and nucleotides which encode these immunoregulatory proteins.
     The proteins, their associated nucleic acids, specific antibodies and
CC
CC
     inhibitors may be used as vaccines for therapeutic or prophylactic
CC
     regulation of an immune response in animals (particularly cats, dogs,
CC
     horses and humans). They may be used to treat autoimmune or infectious
CC
     diseases including allergies, tumours, inflammation and graft rejection,
CC
     and to increase the response from a co-administered antigen. The
CC
     nucleotide sequences can also be used for the recombinant production of a
CC
     protein, while nucleotide fragments are useful as probes, as
CC
     amplification primers and as sources of inhibitory therapeutics (e.g.,
CC
     antisense oligonucleotides). The proteins may be used to raise antibodies
CC
     and to screen for modulators of activity, while the antibodies may be
CC
     used in detection, and in drug targetting
XX
```

```
Sequence 1878 BP; 526 A; 461 C; 434 G; 457 T; 0 U; 0 Other;
SQ
  Query Match
                        87.8%; Score 109.8; DB 3; Length 1878;
  Best Local Similarity
                        97.6%; Pred. No. 8.9e-29;
  Matches 122; Conservative
                               0; Mismatches
                                               2;
                                                   Indels
                                                                 Gaps
                                                                        1;
           2 AAAAACAAAAACCTTTACGTAAC-GTTTTTGCTGGGAGAGAAGACTACGAAGCACATTT 60
Qу
             Db
          93 AAAAAAAAAACCTTTACGTAACTTTTTTTGCTGGGAGAAGACTACGAAGCACATTT 152
Qу
          61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 120
             153 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 212
         121 CTTTG 125
Qy
             IIIII
Db
         213 CTTTG 217
RESULT 6
AAZ55535/c
    AAZ55535 standard; cDNA; 1878 BP.
XX
AC
    AAZ55535;
XX
    14-MAR-2000 (first entry)
DT
XX
DE
    Canine CD154 cDNA complement.
XX
KW
    CD154; CD40 ligand; antibody; canine; feline; inhibitor; immune response;
KW
    immunoregulation; tumour; cancer; autoimmune disease; vaccine; ss.
XX
OS
    Canis familiaris.
XX
FH
    Key
                   Location/Qualifiers
FT
    CDS
                   complement(813. .1595)
FT
                    /*tag= a
FT
                   /product= "Canine CD154"
XX
PN
    WO9961618-A2.
XX
PD
    02-DEC-1999.
XX
PF
                  99WO-US011942.
    28-MAY-1999;
XX
PR
    29-MAY-1998;
                  98US-0087306P.
XX
PA
     (HESK-) HESKA CORP.
XX
PΙ
    Sim G, Yang S, Dreitz MJ, Wonderling RS;
XX
DR
    WPI; 2000-072623/06.
    P-PSDB; AAY58215.
DR
XX
PT
    Nucleic acids encoding immunoregulatory proteins from cats or dogs,
    useful for treating or preventing e.g. tumors or autoimmune disease.
PT
XX
PS
    Claim 1f; Page 209-210; 264pp; English.
XX
CC
    Sequences AAZ55533-Z55539 represent full-length or partial cDNAs encoding
CC
    canine CD154 (CD40 ligand), while sequences AAZ55540-Z55545 represent
```

```
CC
    feline CD154 cDNA sequences. The invention relates to canine interleukin-
     4 (IL-4), canine or feline Flt-3 ligand, canine or feline CD40, canine or
CC
     feline CD154, canine IL-5, canine IL-13, feline interferon-alpha (IFN-
CC
CC
    alpha) and feline granulocyte macrophage colony-stimulating factor
     (GMCSF), and nucleotides which encode these immunoregulatory proteins.
CC
    The proteins, their associated nucleic acids, specific antibodies and
CC
    inhibitors may be used as vaccines for therapeutic or prophylactic
CC
CC
    regulation of an immune response in animals (particularly cats, dogs,
CC
    horses and humans). They may be used to treat autoimmune or infectious
    diseases including allergies, tumours, inflammation and graft rejection,
CC
CC
    and to increase the response from a co-administered antigen. The
CC
    nucleotide sequences can also be used for the recombinant production of a
CC
    protein, while nucleotide fragments are useful as probes, as
CC
    amplification primers and as sources of inhibitory therapeutics (e.g.,
CC
    antisense oligonucleotides). The proteins may be used to raise antibodies
CC
    and to screen for modulators of activity, while the antibodies may be
CC
    used in detection, and in drug targetting
XX
SQ
    Sequence 1878 BP; 457 A; 434 C; 461 G; 526 T; 0 U; 0 Other;
                        87.8%; Score 109.8; DB 3;
                                                   Length 1878;
  Best Local Similarity
                        97.6%; Pred. No. 8.9e-29;
 Matches 122; Conservative
                               0; Mismatches
                                                                        1;
                                                2;
                                                   Indels
Qу
           2 AAAAACAAAAAACCTTTACGTAAC-GTTTTTGCTGGGAGAGAAGACTACGAAGCACATTT 60
             Db
        61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 120
Qу
             1726 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 1667
Db
         121 CTTTG 125
Qу
             ++++
Db
        1666 CTTTG 1662
RESULT 7
AAT05790
ID
    AAT05790 standard; DNA; 620 BP.
XX
AC
    AAT05790;
XX
DT
    28-FEB-1996 (first entry)
XX
DE
    Human CD40 ligand exon 1 chromosomal DNA sequence.
XX
KW
    Exon; intron; chromosome; human CD40 ligand; PCR; amplififaction; primer;
KW
    probe; antisense; gene therapy; HIM syndrome; ds.
XX
OS
    Homo sapiens.
XX
FΗ
                   Location/Qualifiers
    Key
FT
    CDS
                   164. .319
FT
                   /*tag= a
                   /product= "CD40 ligand a.a. 1-51"
FT
FT
                   /note= "exon 1"
\mathbf{FT}
    intron
                   320. .620
FT
                   /*tag= b
FT
                   /cons_splice= 5'site:No, 3'site:Yes
FT
                   /note= "This sequence only shows part of the 5' of the
```

```
FT
                    intron sequence. The 3' splice site of the intron is
                    shown in AAT05791"
FT
XX
     JP07163362-A.
ΡN
XX
     27-JUN-1995.
PD
XX
PF
     25-NOV-1993;
                   93JP-00295102.
XX
PR
     21-OCT-1993;
                   93JP-00263258.
XX
PA
     (MITP ) MITSUBISHI YUKA BCL KK.
XX
DR
     WPI; 1995-260044/34.
DR
     P-PSDB; AAR80834.
XX
     Novel chromosomal DNA encoding a CD40 ligand and probes - useful for
PT
     diagnosis and gene therapy of HIM syndrome.
PT
XX
PS
     Claim 1; Page 5-6; 14pp; Japanese.
XX
     The nucleotide sequence of exon 1 and surrounding 5' untranslated region
CC
     and part of the 3' intron sequence from the chromosomal DNA sequence
CC
CC
     encoding the human CD40 ligand. The gene was isolated using an amplified
     fragment of the gene as a probe. The probe was a fragment amplified by
CC
     the primers AAQ99758-9. The CD40 ligand primary gene is approx. 11 kb
CC
CC
     long and consists of 5 exons and 4 intron sequences (AAT05790-4). The
CC
     sequence of the coding region was used to design a set of
     oligonucleotides (AAQ99760-73) which can be used as antisense probes for
CC.
CC
     use in diagnosis and gene therapy of HIM syndrome
XX
SQ
     Sequence 620 BP; 158 A; 138 C; 133 G; 191 T; 0 U; 0 Other;
  Query Match
                         77.6%; Score 97; DB 2; Length 620;
  Best Local Similarity
                         100.0%; Pred. No. 2.4e-24;
          97; Conservative
                                0; Mismatches
                                                     Indels
Qу
          29 TTTGCTGGGAGAGAGACTACGAAGCACATTTTCCAGGAAGTGTGGGCTGCAACGATTGT 88
             Db
           1 TTTGCTGGGAGAGACTACGAAGCACATTTTCCAGGAAGTGTGGGCTGCAACGATTGT 60
Qу
          89 GCGCTCTTAACTAATCCTGAGTAAGGTGGCCACTTTG 125
             Db
          61 GCGCTCTTAACTAATCCTGAGTAAGGTGGCCACTTTG 97
RESULT 8
AAS46601
     AAS46601 standard; DNA; 7057 BP.
XX
AC
    AAS46601;
XX
DT
     18-DEC-2001 (first entry)
XX
DΕ
     Tumour suppressor gene derived chemically modified sequence #323.
XX
KW
     Human; tumour suppressor gene; oncogene; antitumour; cytostatic; cancer;
KW
     tumour; CpG dinucleotide; single-nucleotide polymorphism; SNP;
KW
     cytosine methylation; ds.
XX
OS
     Homo sapiens.
```

```
XX
PN
    WO200168912-A2.
XX
PD
    20-SEP-2001.
XX
PF
    15-MAR-2001; 2001WO-EP002955.
XX
    15-MAR-2000; 2000DE-01013847.
PR
    06-APR-2000; 2000DE-01019058.
PR
    07-APR-2000; 2000DE-01019173.
PR
    30-JUN-2000; 2000DE-01032529.
PR
    01-SEP-2000; 2000DE-01043826.
PR
XX
PA
     (EPIG-) EPIGENOMICS AG.
XX
PΙ
    Olek A, Piepenbrock C,
                            Berlin K;
XX
DR
    WPI; 2001-602752/68.
XX
    Fragments of chemically modified genes associated with tumor suppressor
PT
PT
    genes and oncogenes, useful in designing primers and probes for analyzing
PT
    diseases associated with cytosine methylation state e.g. cancer.
XX
PS
    Claim 1; SEQ ID NO 323; 27pp; English.
XX
CC
    The invention relates to a nucleic acid comprising a sequence of 18
CC
    bases, of a segment of chemically pretreated DNA (CP DNA) e.g. with
CC
    bisulphite, of genes associated with tumour suppression and oncogenes
CC
    having a sequence taken from 536 (actually 533 since numbers 408, 458 and
    500 are missing from the sequence listing) sequences (Ss) and sequences
CC
CC
    complementary to (Ss). The nucleic acid may be a peptide nucleic acid-
CC
    oligomer (PNA) of at least 9 nucleotides and may form part of a set of
CC
    probes for detecting the cytosine methylation state and/or single
    nucleotide polymorphisms and also to be used in an array for analysing
CC
CC
    diseases associated with CpG dinucleotides e.g. cancers and tumours. The
CC
    probes can also be used in a method for ascertaining genetic and/or
CC
    epigenetic parameters for the diagnosis and/or therapy of existing
CC
    diseases or the predisposition to specific diseases, by analysing
CC
    cytosine methylations. The parameters may be compared to another set of
CC
    genetic and/or epigenetic parameters, the differences serving as basis
CC
    for diagnosis and/or prognosis events which are disadvantageous to
CC
    patients. The present sequence is one of the 533 genomic sequences
CC
    derived from tumour suppressor genes and oncogenes. Note: The sequence
CC
    data for this patent did not form part of the printed specification, but
CC
    was obtained in electronic format directly from WIPO at
CC
    ftp.wipo.int/pub/published_pct sequences
XX
SQ
    Sequence 7057 BP; 2260 A; 51 C; 1534 G; 3212 T; 0 U; 0 Other;
 Query Match
                        74.9%;
                               Score 93.6; DB 4;
                                                  Length 7057;
 Best Local Similarity
                        84.7%; Pred. No. 9.4e-23;
 Matches 105; Conservative
                               0; Mismatches
                                               19;
                                                   Indels
                                                                        0;
                                                             0;
                                                                 Gaps
Qу
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
             Db
        4850 AAAAATAAAAATTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
Qу
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
               Db
```

```
· · Qy
                1111
          4970 TTTG 4973
 Db
 RESULT 9
 ABL33848
 ID
      ABL33848 standard; DNA; 7057 BP.
 XX
 AC
      ABL33848;
 XX
      26-MAR-2002 (first entry)
 DT
 XX
 DE
      Human immune system associated gene SEQ ID NO: 1821.
 XX
 KW
      Human; immune system disease; cytosine methylation; antiasthmatic;
      antiarteriosclerotic; antianaemic; cytostatic; nootropic;
 KW
 KW
      neuroprotective; anti-HIV; anticonvulsant; ophthalmological;
 KW
      antirheumatic; antiarthritic; antidiabetic; antipsoriatic;
      antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;
 KW
      acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;
 KW
      neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;
 KW
 KW
 XX
 os
      Homo sapiens.
 XX
 PN
      WO200200928-A2.
 XX
 PD
      03-JAN-2002.
 XX
 PF
      02-JUL-2001; 2001WO-EP007537.
 XX
 PR
      30-JUN-2000; 2000DE-01032529.
 PR
      01-SEP-2000; 2000DE-01043826.
 XX
 PA
      (EPIG-) EPIGENOMICS AG.
 XX
 PΙ
      Olek A, Piepenbrock C, Berlin K;
 XX
 DR
      WPI; 2002-130909/17.
 XX
      Nucleic acid comprising fragment of chemically modified gene, useful for
 PT
      diagnosis and treatment of diseases associated with abnormal cytosine
 PT
 PΤ
      methylation.
 XX
 PS
      Claim 1; SEQ ID NO 1821; 32pp + Sequence Listing; German.
 XX
 CC
      The present invention provides a number of human immune system associated
 CC
      genes which are modified by the methylation of cytosines. The sequences
      can be used in the diagnosis and treatment of immune system disorders,
 CC
 CC
      including eye diseases such as retinopathy, neovascular glaucoma and
 CC
      macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid
 CC
      leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
 CC
      rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
 CC
      diseases. The present sequence is a gene of the invention
 XX
 SQ
      Sequence 7057 BP; 2260 A; 51 C; 1534 G; 3212 T; 0 U; 0 Other;
                           74.9%; Score 93.6; DB 6;
   Query Match
                                                       Length 7057;
   Best Local Similarity
                           84.7%; Pred. No. 9.4e-23;
   Matches 105; Conservative
                                  0; Mismatches
                                                    19; Indels
                                                                   0; Gaps
                                                                               0;
```

122 TTTG 125

```
2 AAAAACAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61
Qу
              31111 131111 171111111111111111 171111111 1711111 1711111 1
         4850 AAAAATAAAAATTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
Db
Qу
           62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
                Db
         4910 TTAGGAAGTGTGGGTTGTAACGATTGTGCGTTTTTAATTTAATTTTGAGTAAGGTGGTTAT 4969
          122 TTTG 125
Qу
         4970 TTTG 4973
Db
RESULT 10
ABL34594
ID
     ABL34594 standard; DNA; 7057 BP.
XX
AC
     ABL34594;
XX
DT
     26-MAR-2002
                  (first entry)
XX
DE
     Human metastasis associated gene SEQ ID NO: 147.
XX
KW
    Metastasis associated gene; cytostatic; gene therapy; cancer;
KW
     cytosine methylation; gene; ds.
XX
os
     Homo sapiens.
XX
PN
    WO200177376-A2.
XX
PD
     18-OCT-2001.
XX
     06-APR-2001; 2001WO-EP003970.
PF
XX
     06-APR-2000; 2000DE-01019058.
PR
     07-APR-2000; 2000DE-01019173.
PR
     30-JUN-2000; 2000DE-01032529.
PR
PR
     01-SEP-2000; 2000DE-01043826.
XX
PA
     (EPIG-) EPIGENOMICS AG.
XX
ΡI
    Olek A, Piepenbrock C,
                             Berlin K;
XX
    WPI; 2002-010922/01.
DR
XX
PT
    New nucleic acid derived from chemically treated metastasis genes, useful
PT
     for diagnosis of cancers by analysis of cytosine methylation, also for
PT
     treatment.
XX
PS
    Claim 1; SEQ ID NO 147; 23pp + Sequence Listing; English.
XX
CC
    The present invention provides a number of human metastasis associated
CC
     genes which are modified by cytosine methylation. The sequences can be
CC
     used in the diagnosis and treatment of cancer. The present sequence is
CC
    one of the genes of the invention
XX
    Sequence 7057 BP; 2260 A; 51 C; 1534 G; 3212 T; 0 U; 0 Other;
SQ
                         74.9%;
  Query Match
                                 Score 93.6; DB 6;
  Best Local Similarity
                         84.7%; Pred. No. 9.4e-23;
```

```
Matches 105; Conservative
                              0; Mismatches
                                              19;
                                                   Indels
                                                                Gaps
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGGAAGACTACGAAGCACATTTT 61
Qу
             4850 AAAAATAAAAAATTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
Db
Qy
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
               Db
         122 TTTG 125
Qу
             1111
       4970 TTTG 4973
Db
RESULT 11
ABL70407
ID
    ABL70407 standard; DNA; 7057 BP.
XX
AC
    ABL70407;
XX
    01-JUL-2002 (first entry)
DΤ
XX
DE
    Chemically treated cell signalling DNA sequence#149.
XX
KW
    Cell signalling; cytosine methylation; cell signalling disease; cancer;
KW
    tumour; cytostatic; ds.
XX
OS
    Unidentified.
XX
PN
    WO200202807-A2.
XX
PD
    10-JAN-2002.
XX
PF
    29-JUN-2001; 2001WO-EP007471.
XX
PR
    30-JUN-2000; 2000DE-01032529.
    01-SEP-2000; 2000DE-01043826.
XX
PA
    (EPIG-) EPIGENOMICS AG.
XX
PΤ
    Olek A, Piepenbrock C, Berlin K;
XX
DR
    WPI; 2002-154758/20.
XX
PT
    Nucleic acid, useful for diagnosis and therapy of diseases associated
PT
    with cell signaling e.g. cancer, comprises chemically modified genomic
PT
    sequences of genes associated with cell signaling.
XX
PS
    Claim 1; SEQ ID NO 297; 24pp + Sequence Listing; English.
XX
CC
    The invention relates to a nucleic acid comprising a sequence of at least
    18 bases of a segment of chemically pretreated DNA of genes associated
CC
CC
    with cell signalling. The activity of the modified sequences of the
CC
    invention may be described as cytostatic. The object of the invention is
CC
    to provide the chemically modified DNA of genes associated with cell
CC
    signalling, as well as oligonucleotides and/or PNA-oligomers for
CC
    detecting cytosine methylations, as well as a method which is
CC
    particularly suitable for the diagnosis and/or therapy of genetic and
CC
    epigenetic parameters of genes associated with cell signalling. The
    chemically modified DNA provided by the invention is useful for diagnosis
```

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CC
    and therapy of diseases such as solid tumours and cancer. The sequences
    given in records ABL70111-ABL70626 represent chemically pre-treated
CC
CC
    genomic DNA's of genes associated with cell signalling. Note: The
CC
    sequence data for this patent is not represented in the printed
    specification, but is based on sequence information supplied by the
CC
CC
    European Patent Office
XX
    Sequence 7057 BP; 2260 A; 51 C; 1534 G; 3212 T; 0 U; 0 Other;
SO
                        74.9%;
 Ouery Match
                               Score 93.6; DB 6;
                                                 Length 7057;
 Best Local Similarity
                        84.7%;
                                Pred. No. 9.4e-23;
 Matches 105; Conservative
                               0; Mismatches
                                               19;
                                                   Indels
                                                                        0:
                                                                Gaps
Qу
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61 ·
             Qу
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
               11111111111
Db
        4910 TTAGGAAGTGTGGGTTGTAACGATTGTGCGTTTTTAATTTAATTTTGAGTAAGGTGGTTAT 4969
         122 TTTG 125
Qy
             IIIII
Db
        4970 TTTG 4973
RESULT 12
AAS61354
ID
    AAS61354 standard; DNA; 7057 BP.
XX
AC
    AAS61354;
XX
DT
    29-JAN-2002 (first entry)
XX
DE
    Human gene regulation-associated gene oligonucleotide #309.
XX
KW
    Human; Gene regulation-associated gene; severe combined immunodeficiency;
KW
    cardiac damage; inflammatory response; Haemophilia; Werner syndrome;
KW
    asthma; HDR syndrome; congenital heart defect; Saethre-Chotzen syndrome;
KW
    renal disease; Preeclampsia; cardiac allograft vascular disease;
KW
    colorectal cancer; thyroid cancer; oesophageal cancer; ds; tumour;
KW
    immunostimulant; cardiant; antiinflammatory; coagulant; antiasthmatic;
KW
    nephrotropic; gynecological; anti-tumour; immunosuppressive; cytostatic.
XX
os
    Homo sapiens:
XX
PN
    WO200177375-A2.
XX
PD
    18-OCT-2001.
XX
PF
    06-APR-2001; 2001WO-EP003968.
XX
PR
    06-APR-2000; 2000DE-01019058.
    07-APR-2000; 2000DE-01019173.
PR
    30-JUN-2000; 2000DE-01032529.
PR
    01-SEP-2000; 2000DE-01043826.
PR
XX
PA
    (EPIG-) EPIGENOMICS AG.
XX
ΡI
    Olek A,
            Piepenbrock C,
                           Berlin K;
XX
```

```
DR
     WPI; 2002-017470/02.
XX
PT
    New nucleic acid sequences from chemically modified genes associated with
PT
     gene regulation, useful for analyzing cytosine methylations for diagnosis
PT
     and therapy of diseases e.g. severe combined immunodeficiency disease.
XX
PS .
    Disclosure; SEQ ID NO 317; 26pp; English.
XX
CC
    The invention relates to 224 nucleic acid sequences comprising at least
    18 bases of a chemically pretreated gene associated with gene regulation
CC
CC
     selected from 43 known genes (or complementary sequences). The chemical
CC
    pretreatment converts cytosine bases unmethylated at the 5-position to
CC
    uracil or another base with hybridisation behaviour dissimilar to
CC
    cytosine, to enable analysis of cytosine methylations. The DNA sequences,
CC
    oligomers (or sets/arrays) and method are useful in the diagnosis of
    diseases (or predisposition to diseases) associated with gene regulation
CC
CC
    and in therapy of such diseases, by enabling analysis of the cytosine
    methylation patterns of such genes, kits are provided. They are
CC
CG
    especially useful in diagnosis and therapy of e.g. severe combined
CC
    immunodeficiency disease, cardiac disorders, haemophilia, solid tumours
CC
    and cancer, Werner syndrome, asthma, HDR syndrome, Saethre-Chotzen
CC
    syndrome, renal disease, preeclampsia, graft versus-host disease. The
CC
    present sequence is a sequence included in the sequence data for this
CC
    specification and is associated with the human gene regulation-associated
CC
    genes. Note: The sequence data for this patent did not form part of the
CC
    printed specification, but was obtained in electronic format directly
CC
    from WIPO at ftp.wipo.int/pub/published pct sequences
XX
SQ
    Sequence 7057 BP; 2260 A; 51 C; 1534 G; 3212 T; 0 U; 0 Other;
  Query Match
                        74.9%;
                               Score 93.6; DB 6;
                                                  Length 7057;
  Best Local Similarity
                        84.7%; Pred. No. 9.4e-23;
 Matches 105; Conservative
                              0;
                                 Mismatches
                                                  Indels
                                                                       0;
Qу
          Db
        4850 AAAAATAAAAATTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qy
               Db
        122 TTTG 125
Qy
             IIII
Db
        4970 TTTG 4973
RESULT 13
ADS99855
ID
    ADS99855 standard; DNA; 7057 BP.
·ХХ
AC
    ADS99855;
XX
DT
    02-DEC-2004 (first entry)
XX
DΕ
    Bisulphite treated human gene associated with metastasis #74.
XX
    Human; ds; gene; Bisulphite; metastasis; cancer; cytostatic;
KW 
KW
    DNA methylation; matrix-assisted laser desorption/ionisation; MALDI;
KW
    electrospray; mass spectrometry; CpG dinucleotide; solid tumour.
XX
```

```
os
     Homo sapiens.
XX
     US2003148327-A1.
PN
XX
     07-AUG-2003.
PD
XX
     21-JAN-2003; 2003US-00240485.
PF
XX
PR
     06-APR-2000; 2000DE-01019058.
     07-APR-2000; 2000DE-01019173.
PR
PR
     30-JUN-2000; 2000DE-01032529.
     01-SEP-2000; 2000DE-01043826.
PR
     06-APR-2001; 2001WO-EP003970.
PR
XX
PA
     (OLEK/) OLEK A.
PA
     (PIEP/) PIEPENBROCK C.
PA
     (BERL/) BERLIN K.
XX
PΙ
     Olek A, Piepenbrock C,
                              Berlin K;
XX
    WPI; 2002-010922/01.
DR
XX
    New nucleic acid derived from chemically treated metastasis genes, useful
PT
PT
     for diagnosis of cancers by analysis of cytosine methylation, also for
PT
     treatment.
XX
PS
    Claim 1; SEQ ID NO 147; 9pp; English.
XX
     The invention relates to a nucleic acid comprising at least 18 bases from
CC
     a segment of the chemically pretreated DNA of genes associated with
CC
    metastasis, i.e. any of ADS99709-ADS99906 human genomic sequences or any
CC
CC
    of the 19 sequences appearing as ADS99911-ADS99929. SEQ ID 2,4,6 etc are
CC
     the complements of SEQ ID 1,3,5, etc. Also included are an oligomer
CC
     (particularly an oligonucleotide or peptide nucleic acid) comprising at
CC
     least one base sequence of at least 9 bases which hybridises to (or is
CC
     identical with) the sequences referred to above, producing an array of
CC
    the oligomers on a carrier, obtaining genetic and/or epigenetic
CC
    parameters for diagnosis and/or therapy of diseases (or predisposition to
CC
    them) by analysis of cytosine methylation and a kit comprising a
CC
    bisulphite (disulphite or hydrogen sulphite) and the oligomers. In the
CC
    method of above 5-unmethylated cytosines in a genomic DNA sample are
CC
    converted chemically to uracil, or another base with hybridisation
CC
    properties different from those of cytosine, then fragments of the
    treated DNA amplified (particularly by polymerase chain reaction) using
CC.
CC
    the oligomers and a polymerase (preferably heat stable) to produce
CC
    labelled amplicons. These are tested for hybridisation to an array of
CC
    oligomers and any hybridisation detected. The amplicons are labelled with
CC
    fluorescent or radioactive markers, or with a detachable mass marker to
CC
    allow their detection by mass spectrometry, specifically using the matrix
CC
    -assisted laser desorption/ionisation (MALDI) or electrospray techniques.
CC
    To improve detection in the mass spectrometer, fragments formed in the
CC
    instrument have only a single net charge (positive or negative). The
CC
    genomic DNA is from e.g. a cell line, biopsy sample, blood, or paraffin-
CC
    embedded tissue sample. Oligonucleotides or peptide-nucleic acids that
CC
    are complementary to (or identical with) parts of the nuclei acids listed
    above may be used as primers for amplification of the nucleic acids or
CC
CC
    their complements, and for determining cytosine methylation status and/or
CC
    single nucleotide polymorphisms in metastasis-related genes. They can be
CC
    used for analysis of diseases associated with methylation of CpG
CC
    dinucleotides and to determine (epi)genetic parameters for diagnosis
    and/or therapy of disease (or predisposition). The genomic DNA sequences
```

```
CC
    are useful for diagnosis and therapy of solid tumours and cancer. The
CC
    present sequence is a bisulphite treated human gene associated with
    metastasis. Note: The sequence data for this patent did not form part of
CC
    the printed specification, but was obtained in electronic format directly
CC
    from USPTO at seqdata.uspto.gov/sequence.html?DocID=20030148327.
CC
XX
    Sequence 7057 BP; 2260 A; 51 C; 1534 G; 3212 T; 0 U; 0 Other;
SQ
                        74.9%; Score 93.6; DB 7;
  Ouerv Match
                                                 Length 7057;
  Best Local Similarity
                       84.7%; Pred. No. 9.4e-23;
 Matches 105; Conservative
                             0; Mismatches
                                              19;
                                                  Indels
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61
Qy
             4850 AAAAATAAAAATTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
Db
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qy
              Db
        122 TTTG 125
Qу
             1111
        4970 TTTG 4973
Db
RESULT 14
AAS46602/c
    AAS46602 standard; DNA; 7057 BP.
XX
AC
    AAS46602;
XX
DΤ
    18-DEC-2001 (first entry)
XX
DE
    Tumour suppressor gene derived chemically modified sequence #324.
XX
KW
    Human; tumour suppressor gene; oncogene; antitumour; cytostatic; cancer;
KW
    tumour; CpG dinucleotide; single-nucleotide polymorphism; SNP;
KW
    cytosine methylation; ds.
XX
os
    Homo sapiens.
XX
PN
    WO200168912-A2.
XX
PD
    20-SEP-2001.
XX
PF
    15-MAR-2001; 2001WO-EP002955.
XX
PR
    15-MAR-2000; 2000DE-01013847.
PR
    06-APR-2000; 2000DE-01019058.
    07-APR-2000; 2000DE-01019173.
PR
PR
    30-JUN-2000; 2000DE-01032529.
    01-SEP-2000; 2000DE-01043826.
PR
XX
PA
    (EPIG-) EPIGENOMICS AG.
XX
PΙ
    Olek A, Piepenbrock C, Berlin K;
XX
DR
    WPI; 2001-602752/68.
XX
PΤ
    Fragments of chemically modified genes associated with tumor suppressor
    genes and oncogenes, useful in designing primers and probes for analyzing
```

```
PT
    diseases associated with cytosine methylation state e.g. cancer.
XX
PS
    Claim 1; SEQ ID NO 324; 27pp; English.
XX
CC
    The invention relates to a nucleic acid comprising a sequence of 18
    bases, of a segment of chemically pretreated DNA (CP DNA) e.q. with
CC
    bisulphite, of genes associated with tumour suppression and oncogenes
CC
CC
    having a sequence taken from 536 (actually 533 since numbers 408, 458 and
CC
    500 are missing from the sequence listing) sequences (Ss) and sequences
CC
    complementary to (Ss). The nucleic acid may be a peptide nucleic acid-
    oligomer (PNA) of at least 9 nucleotides and may form part of a set of
CC
CC
    probes for detecting the cytosine methylation state and/or single
CC
    nucleotide polymorphisms and also to be used in an array for analysing
CC
    diseases associated with CpG dinucleotides e.g. cancers and tumours. The
CC
    probes can also be used in a method for ascertaining genetic and/or
CC
    epigenetic parameters for the diagnosis and/or therapy of existing
CC
    diseases or the predisposition to specific diseases, by analysing
CC
    cytosine methylations. The parameters may be compared to another set of
CC
    genetic and/or epigenetic parameters, the differences serving as basis
CC
    for diagnosis and/or prognosis events which are disadvantageous to
CC
    patients. The present sequence is one of the 533 genomic sequences
CC
    derived from tumour suppressor genes and oncogenes. Sequences with even
CC
    numbered Seq ID numbers are the complementary sequence of the
CC
    corresponding odd numbered sequence (e.g. ID 2 and ID1, ID 536 and ID
    535, except for those whose partner sequence is missing). Note: The
CC
CC
    sequence data for this patent did not form part of the printed
CC
    specification, but was obtained in electronic format directly from WIPO
CC
    at ftp.wipo.int/pub/published pct sequences
XX
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             Db
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        2088 TTT 2086
Db
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AC
    ABL33849;
XX
    26-MAR-2002 (first entry)
DΤ
XX
DE
    Human immune system associated gene SEQ ID NO: 1822.
XX
KW
    Human; immune system disease; cytosine methylation; antiasthmatic;
KW
    antiarteriosclerotic; antianaemic; cytostatic; nootropic;
    neuroprotective; anti-HIV; anticonvulsant; ophthalmological;
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KW
    antirheumatic; antiarthritic; antidiabetic; antipsoriatic;
    antiinflammatory; cancer; eye disease; arteriosclerosis; anaemia;
KW
    acute myeloid leukaemia; Alzheimer's disease; AIDS; epilepsy;
KW
KW 
    neurofibromatosis; rheumatoid arthritis; psoriasis; bowel disease; gene;
KW
    ds.
XX
os
    Homo sapiens.
XX
PN
    WO200200928-A2.
XX
PD
    03-JAN-2002.
XX
    02-JUL-2001; 2001WO-EP007537.
PF
XX
    30-JUN-2000; 2000DE-01032529.
PR
    01-SEP-2000; 2000DE-01043826.
PR
XX
PΑ
    (EPIG-) EPIGENOMICS AG.
XX
PΙ
    Olek A,
            Piepenbrock C,
                          Berlin K;
XX
DR
    WPI; 2002-130909/17.
XX
PΤ
    Nucleic acid comprising fragment of chemically modified gene, useful for
PT
    diagnosis and treatment of diseases associated with abnormal cytosine
PT
    methylation.
XX
PS
    Claim 1; SEQ ID NO 1822; 32pp + Sequence Listing; German.
ΧX
CC
    The present invention provides a number of human immune system associated
CC
    genes which are modified by the methylation of cytosines. The sequences
CC
    can be used in the diagnosis and treatment of immune system disorders,
CC
    including eye diseases such as retinopathy, neovascular glaucoma and
CC
    macular degeneration, arteriosclerosis, anaemia, cancer, acute myeloid
    leukaemia, Alzheimer's disease, AIDS, epilepsy, neurofibromatosis,
CC
CC
    rheumatoid arthritis, psoriasis and inflammatory/ulcerative bowel
CC
    diseases. The present sequence is a gene of the invention
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 Best Local Similarity
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Qу
            Db
        Qу
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
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Search completed: April 25, 2006, 10:12:51
Job time : 287 secs
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SCORE Search Results Details for Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rni.

Score Home Page Retrieve Application List SCORE System Overview SCORE FAQ Comments / Sugg

This page gives you Search Results detail for the Application 10088319 and Search Result us-10-08 2_copy_331_455.rni.

start

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GenCore version 5.1.7 Copyright (c) 1993 - 2006 Biocceleration Ltd.

OM nucleic - nucleic search, using sw model

Run on: April 25, 2006, 10:13:01; Search time 104 Seconds

(without alignments)

2136.492 Million cell updates/sec

Title: US-10-088-319-2_COPY_331_455

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Gapop 10.0 , Gapext 1.0

Searched: 1303057 seqs, 888780828 residues

Total number of hits satisfying chosen parameters: 2606114

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents NA:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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	2	112	89.6	2395	3	US-09-875-453B-9	Sequence 9, Appli
	3	109.8	87.8	1878	3	US-09-322-409-64	Sequence 64, Appl
С	4	109.8	87.8	1878	3	US-09-322-409-66	Sequence 66, Appl
, •	5	109.8	87.8	1878	3	US-09-451-527-64	Sequence 64, Appl
С	6	109.8	87.8	1878	3	US-09-451-527-66	Sequence 66, Appl
	7		29.6	37	3	US-09-875-453B-15	Sequence 15, Appl
	8	32.8	26.2	1830121	3	US-09-557-884-1	Sequence 1, Appli
	9	32.8		1830121	3	US-09-643-990A-1	Sequence 1, Appli
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ALIGNMENTS

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; Sequence 10, Application US/09909595
; Patent No. 6586245
; GENERAL INFORMATION:
; APPLICANT: C. Frank Bennett
; APPLICANT: Brenda F. Baker
; APPLICANT: Jacqueline Wyatt
; APPLICANT: Scott E. Davis
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TITLE OF INVENTION: ANTISENSE MODULATION OF CD40 LIGAND EXPRESSION
  FILE REFERENCE: RTS-0223
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  CURRENT FILING DATE: 2001-07-18
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   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
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RESULT 2
US-09-875-453B-9
; Sequence 9, Application US/09875453B
; Patent No. 6838556
; GENERAL INFORMATION:
  APPLICANT: Kim, Jungsuh P.
  APPLICANT: Starr, Douglas B.
  APPLICANT: Tam, Albert W.
  APPLICANT: Laurance, Megan E.
APPLICANT: Michelotti, Emil F.
APPLICANT: Velligan, Mark D.
APPLICANT: Latour, Derek R.
  APPLICANT: Thomas, Rita L.
  APPLICANT: Kongpachith, Ana
  APPLICANT: Sheppard, Liana T.
  APPLICANT: Lim, Moon Young
  APPLICANT: Bruice, Thomas W.
  TITLE OF INVENTION: PROMOTERS FOR REGULATED GENE EXPRESSION
  FILE REFERENCE: 54600-8135.US00
  CURRENT APPLICATION NUMBER: US/09/875,453B
  CURRENT FILING DATE: 2001-06-06
  PRIOR APPLICATION NUMBER: US 60/209,549
  PRIOR FILING DATE: 2000-06-06
  NUMBER OF SEQ ID NOS: 246
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; SEQ ID NO 9
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   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-875-453B-9
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Qу
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US-09-322-409-64
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; Patent No. 6471957
; GENERAL INFORMATION:
  APPLICANT: Sim, Gek-Kee
  APPLICANT: Yang, Shumin
  APPLICANT: Dreitz, Matthew J.
  APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
  FILE REFERENCE: IM-2-C1
  CURRENT APPLICATION NUMBER: US/09/322,409
  CURRENT FILING DATE: 1999-05-28
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  EARLIER FILING DATE: 1998-05-29
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   NAME/KEY: CDS
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Db
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Qу
            Db
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US-09-322-409-66/c
; Sequence 66, Application US/09322409
; Patent No. 6471957
; GENERAL INFORMATION:
 APPLICANT: Sim, Gek-Kee
  APPLICANT: Yang, Shumin
  APPLICANT: Dreitz, Matthew J.
  APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
  FILE REFERENCE: IM-2-C1
  CURRENT APPLICATION NUMBER: US/09/322,409
  CURRENT FILING DATE: 1999-05-28
  EARLIER APPLICATION NUMBER: 60/087,306
  EARLIER FILING DATE: 1998-05-29
  NUMBER OF SEQ ID NOS: 154
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   ORGANISM: Canis familiaris
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; Sequence 64, Application US/09451527
; Patent No. 6482403.
; GENERAL INFORMATION:
 APPLICANT: Sim, Gek-Kee
  APPLICANT: Yang, Shumin
  APPLICANT: Dreitz, Matthew J.
  APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
  FILE REFERENCE: IM-2-C2
  CURRENT APPLICATION NUMBER: US/09/451,527
  CURRENT FILING DATE: 1999-12-01
  EARLIER APPLICATION NUMBER: 09/322,409
  EARLIER FILING DATE: 1999-05-28
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; NUMBER OF SEQ ID NOS: 174
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; Patent No. 6482403
; GENERAL INFORMATION:
 APPLICANT: Sim, Gek-Kee
 APPLICANT: Yang, Shumin
 APPLICANT: Dreitz, Matthew J.
 APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
 FILE REFERENCE: IM-2-C2
  CURRENT APPLICATION NUMBER: US/09/451,527
  CURRENT FILING DATE: 1999-12-01
  EARLIER APPLICATION NUMBER: 09/322,409
  EARLIER FILING DATE: 1999-05-28
  EARLIER APPLICATION NUMBER: 60/087,306
 EARLIER FILING DATE: 1998-05-29
 NUMBER OF SEQ ID NOS: 174
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US-09-875-453B-15
; Sequence 15, Application US/09875453B
; Patent No. 6838556
; GENERAL INFORMATION:
  APPLICANT: Kim, Jungsuh P.
  APPLICANT: Starr, Douglas B.
 APPLICANT: Tam, Albert W.
; APPLICANT: Laurance, Megan E.
  APPLICANT: Michelotti, Emil F.
  APPLICANT: Velligan, Mark D.
  APPLICANT: Latour, Derek R.
  APPLICANT: Thomas, Rita L.
  APPLICANT: Kongpachith, Ana
  APPLICANT: Sheppard, Liana T.
  APPLICANT: Lim, Moon Young
  APPLICANT: Bruice, Thomas W.
  TITLE OF INVENTION: PROMOTERS FOR REGULATED GENE EXPRESSION
  FILE REFERENCE: 54600-8135.US00
  CURRENT APPLICATION NUMBER: US/09/875,453B'
  CURRENT FILING DATE: 2001-06-06
; PRIOR APPLICATION NUMBER: US 60/209,549
; PRIOR FILING DATE: 2000-06-06
 NUMBER OF SEQ ID NOS: 246
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 15
   LENGTH: 37
   TYPE: DNA
   ORGANISM: Homo sapiens
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; Patent No. 6506581
   GENERAL INFORMATION:
        APPLICANT: Fleischmann et al.
        TITLE OF INVENTION: The Nucleotide sequence of
                           the Haemophilus influenzae Rd Genome, Fragments
                           Thereof, and Uses Thereof
        NUMBER OF SEQUENCES: 1
        CORRESPONDENCE ADDRESS:
            ADDRESSEE: Human Genome Sciences, Inc.
```

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STREET: 9410 Key West Avenue
            CITY: Rockville
            STATE: MD
            COUNTRY: USA
            ZIP: 20850
        COMPUTER READABLE FORM:
            MEDIUM TYPE: 3 1/2 inch diskette
            COMPUTER: Dell Pentium
            OPERATING SYSTEM: MS DOS v6.22
            SOFTWARE: ASCII Text
        CURRENT APPLICATION DATA:
         APPLICATION NUMBER: US/09/557,884
            FILING DATE: 25-Apr-2000
            CLASSIFICATION:
        PRIOR APPLICATION DATA:
            APPLICATION NUMBER: 08/476,102
            FILING DATE: JUN-5-1995
        ATTORNEY/AGENT INFORMATION:
            NAME: Michelle S. Marks
            REGISTRATION NUMBER: 41,971
            REFERENCE/DOCKET NUMBER: PB186P3
        TELECOMMUNICATION INFORMATION:
            TELEPHONE: 301-309-8504
            TELEFAX: 301-309-8439
   INFORMATION FOR SEQ ID NO: 1:
       SEQUENCE CHARACTERISTICS:
            LENGTH: 1830121 base pairs
            TYPE: nucleic acid
           STRANDEDNESS: double
            TOPOLOGY: linear
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US-09-557-884-1
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US-09-643-990A-1
; Sequence 1, Application US/09643990A
 Patent No. 6528289
   GENERAL INFORMATION:
       APPLICANT: Robert D. Fleischmann
                 Mark D. Adams
                 Owen White
                 Hamilton O. Smith
                 J. Craig Venter
        TITLE OF INVENTION: The Nucleotide sequence of
                          the Haemophilus influenzae Rd Genome, Fragments
                          Thereof, and Uses Thereof
       NUMBER OF SEQUENCES: 1
       CORRESPONDENCE ADDRESS:
```

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ADDRESSEE: Human Genome Sciences, Inc.
             STREET: 9410 Key West Avenue
             CITY: Rockville,
             STATE: MD
           COUNTRY: USA
             ZIP: 20850
        COMPUTER READABLE FORM:
             MEDIUM TYPE: 3 1/2 inch diskette
             COMPUTER: Dell Pentium
             OPERATING SYSTEM: MS DOS v6.22
             SOFTWARE: ASCII Text
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             APPLICATION NUMBER: US/09/643,990A
             FILING DATE: 23-Aug-2000
             CLASSIFICATION:
        PRIOR APPLICATION DATA:
             APPLICATION NUMBER: 08/487,429
             FILING DATE: 1995-06-07
             APPLICATION NUMBER: 08/426,787
             FILING DATE: 1995-04-21
        ATTORNEY/AGENT INFORMATION:
             NAME: Kenley K. Hoover
             REGISTRATION NUMBER: 40,302
             REFERENCE/DOCKET NUMBER: PB186P1C1
        TELECOMMUNICATION INFORMATION:
             TELEPHONE: 301-610-5790
             TELEFAX: 310-309-8439
   INFORMATION FOR SEQ ID NO: 1:
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; Sequence 1, Application US/10158865
; Patent No. 6846651
; GENERAL INFORMATION:
; APPLICANT: Fleischmann et al.
  TITLE OF INVENTION: Nucleotide Sequence of the Haemophilus Influenzae Rd Genome, Fr
; Patent No. 6846651
; TITLE OF INVENTION: Thereof, and Uses Thereof
; FILE REFERENCE: PB186P2C1D1
; CURRENT APPLICATION NUMBER: US/10/158,865
; CURRENT FILING DATE: 2002-06-03
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; PRIOR FILING DATE: 1995-04-21
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Qу
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RESULT 11
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; Sequence 14179, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
 TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
; TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
; FILE REFERENCE: CL001307
 CURRENT APPLICATION NUMBER: US/09/949,016
 CURRENT FILING DATE: 2000-04-14
 PRIOR APPLICATION NUMBER: 60/241,755
 PRIOR FILING DATE: 2000-10-20
 PRIOR APPLICATION NUMBER: 60/237,768
 PRIOR FILING DATE: 2000-10-03
 PRIOR APPLICATION NUMBER: 60/231,498
 PRIOR FILING DATE: 2000-09-08
  NUMBER OF SEQ ID NOS: 207012
 SOFTWARE: FastSEQ for Windows Version 4.0
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   TYPE: DNA
  ORGANISM: Human
   FEATURE:
   NAME/KEY: misc feature
   LOCATION: (1)...(340380)
   OTHER INFORMATION: n = A, T, C or G
US-09-949-016-14179
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; Sequence 1443, Application US/09248796A
; Patent No. 6747137
; GENERAL INFORMATION:
  APPLICANT: Keith Weinstock et al
  TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO CANDIDA ALBIC
  TITLE OF INVENTION: FOR DIAGNOSTICS AND THERAPEUTICS
  FILE REFERENCE: 107196.132
  CURRENT APPLICATION NUMBER: US/09/248,796A
  CURRENT FILING DATE: 1999-02-12
  PRIOR APPLICATION NUMBER: US 60/074,725
  PRIOR FILING DATE: 1998-02-13
 PRIOR APPLICATION NUMBER: US 60/096,409
 PRIOR FILING DATE: 1998-08-13
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          83 GATTGTGCGCTCTTAACTAATCC 105
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RESULT 13
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; Sequence 15321, Application US/09949016
; Patent No. 6812339
; GENERAL INFORMATION:
  APPLICANT: VENTER, J. Craig et al.
  TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
  TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
  FILE REFERENCE: CL001307
  CURRENT APPLICATION NUMBER: US/09/949,016
  CURRENT FILING DATE: 2000-04-14
  PRIOR APPLICATION NUMBER: 60/241,755
  PRIOR FILING DATE: 2000-10-20
  PRIOR APPLICATION NUMBER: 60/237,768
  PRIOR FILING DATE: 2000-10-03
  PRIOR APPLICATION NUMBER: 60/231,498
  PRIOR FILING DATE: 2000-09-08
  NUMBER OF SEQ ID NOS: 207012
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 15321
   LENGTH: 74790
   TYPE: DNA
   ORGANISM: Human
US-09-949-016-15321
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US-09-107-532A-3181
; Sequence 3181, Application US/09107532A
 Patent No. 6583275
   GENERAL INFORMATION:
        APPLICANT: Lynn A Doucette-Stamm and David Bush
        TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO
                           ENTEROCOCCUS FAECIUM FOR DIAGNOSTICS AND THERAPEUTICS
        NUMBER OF SEQUENCES: 7310
        CORRESPONDENCE ADDRESS:
             ADDRESSEE: GENOME THERAPEUTICS CORPORATION
             STREET: 100 Beaver Street
             CITY: Waltham
             STATE: Massachusetts
             COUNTRY: USA
             ZIP: 02354
        COMPUTER READABLE FORM:
             MEDIUM TYPE: CD/ROM ISO9660
             COMPUTER: PC
             OPERATING SYSTEM:
             SOFTWARE: ASCII
        CURRENT APPLICATION DATA:
             APPLICATION NUMBER: US/09/107,532A
             FILING DATE: 30-Jun-1998
        PRIOR APPLICATION DATA:
             APPLICATION NUMBER: 60/085,598
             FILING DATE: 14 May 1998
             APPLICATION NUMBER: 60/051571
             FILING DATE: July 2, 1997
        ATTORNEY/AGENT INFORMATION:
             NAME: Ariniello, Pamela Deneke
             REGISTRATION NUMBER: 40,489
             REFERENCE/DOCKET NUMBER: GTC-012
        TELECOMMUNICATION INFORMATION:
             TELEPHONE: (781)893-5007
             TELEFAX: (781)893-8277
   INFORMATION FOR SEQ ID NO: 3181:
        SEQUENCE CHARACTERISTICS:
             LENGTH: 1746 base pairs
             TYPE: nucleic acid
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             TOPOLOGY: circular
        MOLECULE TYPE: DNA (genomic)
        HYPOTHETICAL: NO
        ANTI-SENSE: NO
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ORGANISM: Enterococcus faecium

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; Patent No. 6812339
; GENERAL INFORMATION:
; APPLICANT: VENTER, J. Craig et al.
 TITLE OF INVENTION: POLYMORPHISMS IN KNOWN GENES ASSOCIATED
 TITLE OF INVENTION: WITH HUMAN DISEASE, METHODS OF DETECTION AND USES THEREOF
 FILE REFERENCE: CL001307
  CURRENT APPLICATION NUMBER: US/09/949,016
  CURRENT FILING DATE: 2000-04-14
 PRIOR APPLICATION NUMBER: 60/241,755
  PRIOR FILING DATE: 2000-10-20
  PRIOR APPLICATION NUMBER: 60/237,768
  PRIOR FILING DATE: 2000-10-03
  PRIOR APPLICATION NUMBER: 60/231,498
  PRIOR FILING DATE: 2000-09-08
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Qy
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Db
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Search completed: April 25, 2006, 10:53:51

Job time : 107 secs

SCORE 1.3 BuildDate: 12/06/2005

SCORE Search Results Details for Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rnpbm.

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OM nucleic - nucleic search, using sw model

Run on: April 25, 2006, 10:52:12; Search time 430 Seconds

(without alignments)

2403.889 Million cell updates/sec

Title: US-10-088-319-2 COPY 331 455

Perfect score: 125

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Gapop 10.0 , Gapext 1.0

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Post-processing: Minimum Match 0%

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Maximum Match 100%

Listing first 45 summaries

Database : Published Applications NA Main:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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No.		Score		Length		ID 	Description
	1	112	89.6	2395	3	US-09-875-453-9	Sequence 9, Appli
	2	112	89.6	2395	3	US-09-909-595-10	Sequence 10, Appl
	3	112	89.6	2395	8	US-10-484-007-10	Sequence 10, Appl
	4	109.8	87.8	1878	5	US-10-218-654-64	Sequence 64, Appl
С	5	109.8	87.8	1878	5	US-10-218-654-66	Sequence 66, Appl
	6	109.8	87.8	1878	6	US-10-262-439-64	Sequence 64, Appl
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	8	93.6	74.9	7057	6	US-10-311-455-1821	Sequence 1821, Ap
	9	93.6	74.9	7057	6	US-10-240-485-147	Sequence 147, App
	10	93.6	74.9	7057	7	US-10-221-613-317	Sequence 317, App
	11	93.6	74.9	7057	7	US-10-221-714A-323	Sequence 323, App
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С	15	84.6	67.7	7057	7	US-10-221-714A-324	Sequence 324, App
	16	37	29.6	37	3	US-09-875-453-15	Sequence 15, Appl
	17	32.8	26.2	1830121	7	US-10-329-670-1	Sequence 1, Appli
	18	32.8	26.2	1830121	8	US-10-158-865-1	Sequence 1, Appli
	19	32.8	26.2	1830121	9	US-10-981-687-1	Sequence 1, Appli
	20	31.4	25.1	290040	8	US-10-850-591-3	Sequence 3, Appli
	21	31.4	25.1	290040	8	US-10-850-586-3	Sequence 3, Appli
С	22	30.8	24.6	1493	8	US-10-739-930-436	Sequence 436, App
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С	36	29.6	23.7	2144	5	US-10-027-632-100413	Sequence 100413,
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С	38	29.6	23.7	2144	6	US-10-027-632-100413	Sequence 100413,
С	39	29.6	23.7	2144	6	US-10-027-632-103092	Sequence 103092,
	40	29.6	23.7	2150	5	US-10-027-632-97497	Sequence 97497, A
	41	29.6	23.7	2150	6	US-10-027-632-97497	Sequence 97497, A
	42	29.2	23.4	474	. 3	US-09-864-761-11535	Sequence 11535, A
	43	28.8	23.0	500	5	US-10-027-632-104434	Sequence 104434,
	44	28.8	23.0	500	5	US-10-027-632-325533	Sequence 325533,
	45	28.8	23.0	500	6	US-10-027-632-104434	Sequence 104434,

ALIGNMENTS

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RESULT 1
US-09-875-453-9
; Sequence 9, Application US/09875453
; Publication No. US20030027320A1
; GENERAL INFORMATION:
; APPLICANT: Kim, Jungsuh P.
; APPLICANT: Starr, Douglas B.
; APPLICANT: Tam, Albert W.
```

```
APPLICANT: Laurance, Megan E.
  APPLICANT: Michelotti, Emil F.
  APPLICANT: Velligan, Mark D.
  APPLICANT: Latour, Derek R.
  APPLICANT: Thomas, Rita L.
  APPLICANT: Kongpachith, Ana
  APPLICANT: Sheppard, Liana T.
  APPLICANT: Lim, Moon Young
  APPLICANT: Bruice, Thomas W.
  TITLE OF INVENTION: PROMOTERS FOR REGULATED GENE EXPRESSION
  FILE REFERENCE: 4600-0135.30
  CURRENT APPLICATION NUMBER: US/09/875,453
  CURRENT FILING DATE: 2001-06-06
  PRIOR APPLICATION NUMBER: US 60/209,549
  PRIOR FILING DATE: 2000-06-06
  NUMBER OF SEQ ID NOS: 78
  SOFTWARE: FastSEQ for Windows Version 4.0
 SEQ ID NO 9
   LENGTH: 2395
   TYPE: DNA
   ORGANISM: Homo sapiens
US-09-875-453-9
 Query Match
                        89.6%;
                               Score 112; DB 3; Length 2395;
                               Pred. No. 4.7e-29;
 Best Local Similarity
                      99.2%;
 Matches 123; Conservative
                              0; Mismatches
                                               0;
                                                   Indels
                                                                        1:
Qу
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
             Db
        1750 AAAAACAAAAACCTTTACGTAACG-TTTTGCTGGGAGAGAGACTACGAAGCACATTTT 1808
Qy
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
             1809 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 1868
Db
         122 TTTG 125
Qy
             1111
        1869 TTTG 1872
Db
RESULT 2
US-09-909-595-10
; Sequence 10, Application US/09909595
; Publication No. US20030083278A1
 GENERAL INFORMATION:
  APPLICANT: C. Frank Bennett
  APPLICANT: Brenda F. Baker
  APPLICANT: Jacqueline Wyatt
  APPLICANT: Scott E. Davis
  TITLE OF INVENTION: ANTISENSE MODULATION OF CD40 LIGAND EXPRESSION
  FILE REFERENCE: RTS-0223
  CURRENT APPLICATION NUMBER: US/09/909,595
  CURRENT FILING DATE: 2001-07-18
  NUMBER OF SEQ ID NOS: 91
 SEQ ID NO 10
   LENGTH: 2395
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1939)...(2094)
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Query Match
                      89.6%; Score 112; DB 3; Length 2395;
 Best Local Similarity 99.2%; Pred. No. 4.7e-29;
 Matches 123; Conservative
                           0; Mismatches
                                         0;
                                              Indels
                                                                 1;
Qy
          2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61
            Db
       1750 AAAAACAAAAAACCTTTACGTAACG-TTTTGCTGGGAGAGAGACTACGAAGCACATTTT 1808
         62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qу
            1809 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 1868
        122 TTTG 125
Qy
            IIII
       1869 TTTG 1872
Db
RESULT 3
US-10-484-007-10
; Sequence 10, Application US/10484007
; Publication No. US20040259824A1
; GENERAL INFORMATION:
  APPLICANT: C. Frank Bennett
  APPLICANT: Brenda F. Baker
  APPLICANT: Jacqueline Wyatt APPLICANT: Scott E. Davis
 APPLICANT: Isis Pharmaceuticals, Inc.
  TITLE OF INVENTION: ANTISENSE MODULATION OF CD40 LIGAND EXPRESSION
 FILE REFERENCE: RTSP-0397
 CURRENT APPLICATION NUMBER: US/10/484,007
  CURRENT FILING DATE: 2004-01-15
 PRIOR APPLICATION NUMBER: 09/909,595
 PRIOR FILING DATE: 2001-07-18
 NUMBER OF SEQ ID NOS: 91
 SEQ ID NO 10
   LENGTH: 2395
   TYPE: DNA
   ORGANISM: Homo sapiens
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (1939)...(2094)
US-10-484-007-10
 Query Match
                     89.6%; Score 112; DB 8; Length 2395;
 Best Local Similarity 99.2%; Pred. No. 4.7e-29;
 Matches 123; Conservative
                           0; Mismatches
                                              Indels
          Qу
            Db
       1750 AAAAACAAAAACCTTTACGTAACG-TTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 1808
Qy
         62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
            Db
       1809 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 1868
        122 TTTG 125
Qу
           +111
Db
       1869 TTTG 1872
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RESULT 4
US-10-218-654-64
; Sequence 64, Application US/10218654
; Publication No. US20030099609A1
; GENERAL INFORMATION:
  APPLICANT: Sim, Gek-Kee
  APPLICANT: Yang, Shumin
  APPLICANT: Dreitz, Matthew J.
  APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
  FILE REFERENCE: IM-2-C1
  CURRENT APPLICATION NUMBER: US/10/218,654
  CURRENT FILING DATE: 2002-08-13
  PRIOR APPLICATION NUMBER: US/09/322,409
  PRIOR FILING DATE: 1999-05-28
  PRIOR APPLICATION NUMBER: 60/087,306
  PRIOR FILING DATE: 1998-05-29
  NUMBER OF SEQ ID NOS: 154
  SOFTWARE: PatentIn Ver. 2.0
 SEQ ID NO 64
   LENGTH: 1878
   TYPE: DNA
   ORGANISM: Canis familiaris
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (284)..(1063)
US-10-218-654-64
 Query Match
                        87.8%; Score 109.8; DB 5;
                                                   Length 1878;
 Best Local Similarity
                        97.6%; Pred. No. 2.6e-28;
 Matches 122; Conservative
                              0; Mismatches
                                                   Indels
                                                                 Gaps
Qy
           2 AAAAACAAAAACCTTTACGTAAC-GTTTTTGCTGGGAGAAGACTACGAAGCACATTT 60
             Db
          93 AAAAAAAAAAACCTTTACGTAACTTTTTTTGCTGGGAGAGAAGACTACGAAGCACATTT 152
          61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 120
Qy
             Db
         153 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 212
         121 CTTTG 125
Qy
             11111
Db
         213 CTTTG 217
RESULT 5
US-10-218-654-66/c
; Sequence 66, Application US/10218654
; Publication No. US20030099609A1
; GENERAL INFORMATION:
  APPLICANT: Sim, Gek-Kee
  APPLICANT:
             Yang, Shumin
  APPLICANT:
             Dreitz, Matthew J.
  APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
  FILE REFERENCE: IM-2-C1
  CURRENT APPLICATION NUMBER: US/10/218,654
  CURRENT FILING DATE: 2002-08-13
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PRIOR APPLICATION NUMBER: US/09/322,409
  PRIOR FILING DATE: 1999-05-28
 PRIOR APPLICATION NUMBER: 60/087,306
 PRIOR FILING DATE: 1998-05-29
 NUMBER OF SEQ ID NOS: 154
  SOFTWARE: PatentIn Ver. 2.0
 SEQ ID NO 66
   LENGTH: 1878
   TYPE: DNA
   ORGANISM: Canis familiaris
US-10-218-654-66
                       87.8%; Score 109.8; DB 5; Length 1878;
 Query Match
 Best Local Similarity 97.6%; Pred. No. 2.6e-28;
 Matches 122; Conservative
                             0; Mismatches
                                                Indels
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Qу
            Dh
       61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 120
Qу
            Db
       1726 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 1667
        121 CTTTG 125
Qу
            11111
       1666 CTTTG 1662
RESULT 6
US-10-262-439-64
; Sequence 64, Application US/10262439
; Publication No. US20030143196A1
; GENERAL INFORMATION:
 APPLICANT: Sim, Gek-Kee
 APPLICANT: Yang, Shumin
  APPLICANT: Dreitz, Matthew J.
  APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
  FILE REFERENCE: IM-2-C2
 CURRENT APPLICATION NUMBER: US/10/262,439
 CURRENT FILING DATE: 2002-09-30
 PRIOR APPLICATION NUMBER: US/09/451,527
 PRIOR FILING DATE: 1999-12-01
 PRIOR APPLICATION NUMBER: 09/322,409
 PRIOR FILING DATE: 1999-05-28
 PRIOR APPLICATION NUMBER: 60/087,306
  PRIOR FILING DATE: 1998-05-29
 NUMBER OF SEQ ID NOS: 174
  SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 64
  LENGTH: 1878
   TYPE: DNA
  ORGANISM: Canis familiaris
   FEATURE:
   NAME/KEY: CDS
   LOCATION: (284)..(1063)
US-10-262-439-64
                       87.8%; Score 109.8; DB 6; Length 1878;
 Query Match
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Best Local Similarity
                       97.6%; Pred. No. 2.6e-28;
  Matches 122; Conservative
                             0; Mismatches
                                             2;
                                                 Indels
                                                             Gaps
                                                                    1;
          2 AAAAACAAAAAACCTTTACGTAAC-GTTTTTGCTGGGAGAGAAGACTACGAAGCACATTT 60
Qу
            93 AAAAAAAAAAACCTTTACGTAACTTTTTTTGCTGGGAGAGAAGACTACGAAGCACATTT 152
Db
          61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 120
Qy
            Db
         153 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 212
         121 CTTTG 125
Qy
            Db
         213 CTTTG 217
RESULT 7
US-10-262-439-66/c
; Sequence 66, Application US/10262439
; Publication No. US20030143196A1
; GENERAL INFORMATION:
  APPLICANT: Sim, Gek-Kee
  APPLICANT:
            Yang, Shumin
  APPLICANT: Dreitz, Matthew J.
  APPLICANT: Wonderling, Ramani S.
  TITLE OF INVENTION: CANINE AND FELINE IMMUNOREGULATORY PROTEINS, NUCLEIC
  TITLE OF INVENTION: ACID MOLECULES, AND USES THEREOF
  FILE REFERENCE: IM-2-C2
  CURRENT APPLICATION NUMBER: US/10/262,439
  CURRENT FILING DATE: 2002-09-30
  PRIOR APPLICATION NUMBER: US/09/451,527
  PRIOR FILING DATE: 1999-12-01
  PRIOR APPLICATION NUMBER: 09/322,409
  PRIOR FILING DATE: 1999-05-28
  PRIOR APPLICATION NUMBER: 60/087,306
  PRIOR FILING DATE: 1998-05-29
  NUMBER OF SEQ ID NOS: 174
  SOFTWARE: PatentIn Ver. 2.0
 SEQ ID NO 66
   LENGTH: 1878
   TYPE: DNA
   ORGANISM: Canis familiaris
US-10-262-439-66
 Query Match
                       87.8%;
                              Score 109.8; DB 6;
                                                Length 1878;
 Best Local Similarity
                       97.6%;
                            Pred. No. 2.6e-28;
 Matches 122; Conservative
                             0; Mismatches
                                                Indels
Qy
          2 AAAAACAAAAAACCTTTACGTAAC-GTTTTTGCTGGGAGAGAAGACTACGAAGCACATTT 60
            1786 AAAAAAAAAACCTTTACGTAACTTTTTTTGCTGGGAGGAGAGACTACGAAGCACATTT 1727
Db
         61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 120
Qу
            Db
        1726 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCA 1667
        121 CTTTG 125
Qy
            1111
Db
        1666 CTTTG 1662
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RESULT 8
US-10-311-455-1821
; Sequence 1821, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
   APPLICANT: OLEK, Alexander
   APPLICANT: PIEPENBROCK, Christian
   APPLICANT: BERLIN, Kurt
   TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Dete
   TITLE OF INVENTION: cytosine methylation
   FILE REFERENCE: 5013.1014
   CURRENT APPLICATION NUMBER: US/10/311,455
   CURRENT FILING DATE: 2002-12-16
   PRIOR APPLICATION NUMBER: PCT/EP01/07537
   PRIOR FILING DATE: 2001-07-02
   PRIOR APPLICATION NUMBER: DE 10032529.7
   PRIOR FILING DATE: 2000-06-30
   PRIOR APPLICATION NUMBER: DE 10043826.1
   PRIOR FILING DATE: 2000-09-01
   NUMBER OF SEQ ID NOS: 2424
 SEQ ID NO 1821
    LENGTH: 7057
    TYPE: DNA
    ORGANISM: Artificial Sequence
    FEATURE:
    OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-311-455-1821
                        74.9%; Score 93.6; DB 6;
  Query Match
                                                 Length 7057;
  Best Local Similarity
                        84.7%; Pred. No. 2.5e-22;
  Matches 105; Conservative
                              0; Mismatches
                                              19;
                                                                      0;
                                                  Indels
                                                               Gaps
Qу
           2 AAAAACAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
             Db
        4850 AAAAATAAAAATTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qу
               Db
Qу
         122 TTTG 125
             1111
Db
        4970 TTTG 4973
RESULT 9
US-10-240-485-147
; Sequence 147, Application US/10240485
; Publication No. US20030148327A1
; GENERAL INFORMATION:
  APPLICANT: OLEK, Alexander
   APPLICANT: PIEPENBROCK, Christian
  APPLICANT: BERLIN, Kurt
   TITLE OF INVENTION: Diagnosis of Diseases Associated with
   TITLE OF INVENTION: Metastasis
   FILE REFERENCE: 5013.1007
  CURRENT APPLICATION NUMBER: US/10/240,485
   CURRENT FILING DATE: 2002-10-02
   PRIOR APPLICATION NUMBER: PCT/EP01/03970
   PRIOR FILING DATE: 2001-04-06
   PRIOR APPLICATION NUMBER: DE 10019058.8
```

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PRIOR FILING DATE: 2000-04-06
  PRIOR APPLICATION NUMBER: DE 10019173.8
  PRIOR FILING DATE: 2000-04-07
  PRIOR APPLICATION NUMBER: DE 10032529.7
 PRIOR FILING DATE: 2000-06-30
 PRIOR APPLICATION NUMBER: DE 10043826.1
 PRIOR FILING DATE: 2000-09-01
 NUMBER OF SEQ ID NOS: 202
 SEO ID NO 147
   LENGTH: 7057
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-240-485-147
                      74.9%; Score 93.6; DB 6; Length 7057;
 Query Match
 Best Local Similarity 84.7%; Pred. No. 2.5e-22;
 Matches 105; Conservative 0; Mismatches 19;
                                                Indels
                                                         0: Gaps
Qу
          2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
            Db
        4850 AAAAATAAAAAATTTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
Qу
         62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
              Db
        122 TTTG 125
Qу
           4970 TTTG 4973
Db
RESULT 10
US-10-221-613-317
; Sequence 317, Application US/10221613
; Publication No. US20040029123A1
; GENERAL INFORMATION:
 APPLICANT: OLEK, ALexander
  APPLICANT: PIEPENBROCK, Christian
  APPLICANT: BERLIN, Kurt
  TITLE OF INVENTION: Diagnosis of Diseases Associated with Cell Cycle
  FILE REFERENCE: 5013.1004
 CURRENT APPLICATION NUMBER: US/10/221,613
 CURRENT FILING DATE: 2002-09-13
  PRIOR APPLICATION NUMBER: PCT/EP01/02945
    DE 10013847.00
    DE 10019058.8
    DE 10019173.8
    DE 10032529.7
   DE 10043826.1
  PRIOR FILING DATE: 2001-03-15
    2000-03-15
    2000-04-06
    2000-04-07
    2000-06-30
    2000-09-01
 NUMBER OF SEQ ID NOS: 428
; SEQ ID NO 317
   LENGTH: 7057
   TYPE: DNA
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ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-221-613-317
                        74.9%;
  Query Match
                               Score 93.6; DB 7;
                                                 Length 7057;
  Best Local Similarity
                        84.7%;
                               Pred. No. 2.5e-22;
 Matches 105; Conservative
                              0; Mismatches
                                                  Indels
                                                               Gaps
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61
Qу
             Db
        4850 AAAAATAAAAATTTTTACGTAACGTTTTTGTTGGGAGAAGATTACGAAGTATATTTT 4909
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qy
               Db
         122 TTTG 125
Qy
             1111
Db
        4970 TTTG 4973
RESULT 11
US-10-221-714A-323
; Sequence 323, Application US/10221714A
; Publication No. US20040048254A1
; GENERAL INFORMATION:
  APPLICANT: OLEK, Alexander
  APPLICANT: PIEPENBROCK, Christian
  APPLICANT: BERLIN, Kurt
  TITLE OF INVENTION: Diagnosis of Diseases Associated with
  TITLE OF INVENTION: tumor suppressor genes and oncogenes
  FILE REFERENCE: 5013.1005
  CURRENT APPLICATION NUMBER: US/10/221,714A
  CURRENT FILING DATE: 2003-01-21
  PRIOR APPLICATION NUMBER: PCT/EP01/02955
  PRIOR FILING DATE: 2001-03-15
  PRIOR APPLICATION NUMBER: DE 10013847.0
  PRIOR FILING DATE: 2000-03-15
  PRIOR APPLICATION NUMBER: DE 10019058.8
  PRIOR FILING DATE: 2000-04-06
  PRIOR APPLICATION NUMBER: DE 10019173.8
  PRIOR FILING DATE: 2000-04-07
  PRIOR APPLICATION NUMBER: DE 10032529.7
  PRIOR FILING DATE: 2000-06-30
  PRIOR APPLICATION NUMBER: DE 10043826.1
  PRIOR FILING DATE: 2000-09-01
  NUMBER OF SEQ ID NOS: 540
 SEQ ID NO 323
   LENGTH: 7057
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-221-714A-323
 Query Match
                       74.9%;
                               Score 93.6; DB 7;
                                                 Length 7057;
 Best Local Similarity
                       84.7%;
                               Pred. No. 2.5e-22;
 Matches 105; Conservative
                              0;
                                 Mismatches
                                             19;
                                                  Indels
Qy
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
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4850 AAAAATAAAAATTTTTACGTAACGTTTTTGTTGGGAGAGAAGATTACGAAGTATATTTT 4909
Db
Qy
         62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
             Db
       122 TTTG 125
Qу
           IIII
Db
       4970 TTTG 4973
RESULT 12
US-10-311-455-1822/c
; Sequence 1822, Application US/10311455
; Publication No. US20030143606A1
; GENERAL INFORMATION:
  APPLICANT: OLEK, Alexander
  APPLICANT: PIEPENBROCK, Christian
  APPLICANT: BERLIN, Kurt
  TITLE OF INVENTION: Diagnosis of Diseases Associated with the Immune System by Dete
  TITLE OF INVENTION: cytosine methylation
  FILE REFERENCE: 5013.1014
  CURRENT APPLICATION NUMBER: US/10/311,455
  CURRENT FILING DATE: 2002-12-16
  PRIOR APPLICATION NUMBER: PCT/EP01/07537
  PRIOR FILING DATE: 2001-07-02
  PRIOR APPLICATION NUMBER: DE 10032529.7
  PRIOR FILING DATE: 2000-06-30
  PRIOR APPLICATION NUMBER: DE 10043826.1
  PRIOR FILING DATE: 2000-09-01
  NUMBER OF SEQ ID NOS: 2424
 SEQ ID NO 1822
   LENGTH: 7057
   TYPE: DNA
   ORGANISM: Artificial Sequence
   OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-311-455-1822
 Query Match
                     67.7%;
                           Score 84.6; DB 6;
                                           Length 7057;
 Best Local Similarity
                    80.5%;
                           Pred. No. 4e-19;
         99; Conservative
 Matches
                          0; Mismatches
                                        24;
                                            Indels
                                                              0;
Qу
         2 AAAAACAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGAACTACGAAGCACATTTT 61
           Db
         62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qу
                      11 1111111 | 111111111111111 | 111 | 1111
       Db
       122 TTT 124
Qy
           IIII
Db
       2088 TTT 2086
RESULT 13
US-10-240-485-148/c
; Sequence 148, Application US/10240485
; Publication No. US20030148327A1
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; GENERAL INFORMATION:
  APPLICANT: OLEK, Alexander
  APPLICANT: PIEPENBROCK, Christian
  APPLICANT: BERLIN, Kurt
  TITLE OF INVENTION: Diagnosis of Diseases Associated with
  TITLE OF INVENTION: Metastasis
  FILE REFERENCE: 5013.1007
  CURRENT APPLICATION NUMBER: US/10/240,485
  CURRENT FILING DATE: 2002-10-02
  PRIOR APPLICATION NUMBER: PCT/EP01/03970
  PRIOR FILING DATE: 2001-04-06
  PRIOR APPLICATION NUMBER: DE 10019058.8
  PRIOR FILING DATE: 2000-04-06
  PRIOR APPLICATION NUMBER: DE 10019173.8
  PRIOR FILING DATE: 2000-04-07
  PRIOR APPLICATION NUMBER: DE 10032529.7
  PRIOR FILING DATE: 2000-06-30
  PRIOR APPLICATION NUMBER: DE 10043826.1
  PRIOR FILING DATE: 2000-09-01
  NUMBER OF SEQ ID NOS: 202
 SEO ID NO 148
   LENGTH: 7057
   TYPE: DNA
   ORGANISM: Artificial Sequence
   FEATURE:
   OTHER INFORMATION: chemically treated genomic DNA (Homo sapiens)
US-10-240-485-148
                       67.7%;
 Query Match
                             Score 84.6; DB 6;
                                               Length 7057;
 Best Local Similarity
                      80.5%;
                              Pred. No. 4e-19;
         99; Conservative
                             0; Mismatches
                                                                    0;
                                            24;
                                                Indels
                                                             Gaps
Qy
          2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACACACGAAGCACATTTT 61
            1 1 11 111111 1111
       Db
         62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qy
                Db
       122 TTT 124
Qу
            111
       2088 TTT 2086
RESULT 14
US-10-221-613-318/c
; Sequence 318, Application US/10221613
; Publication No. US20040029123A1
; GENERAL INFORMATION:
  APPLICANT: OLEK, ALexander
  APPLICANT: PIEPENBROCK, Christian
  APPLICANT: BERLIN, Kurt
  TITLE OF INVENTION: Diagnosis of Diseases Associated with Cell Cycle
  FILE REFERENCE: 5013.1004
  CURRENT APPLICATION NUMBER: US/10/221,613
  CURRENT FILING DATE: 2002-09-13
  PRIOR APPLICATION NUMBER: PCT/EP01/02945
    DE 10013847.00
    DE 10019058.8
    DE 10019173.8
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    2000-06-30
    2000-09-01
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; Publication No. US20040048254A1
; GENERAL INFORMATION:
 APPLICANT: OLEK, Alexander
 APPLICANT: PIEPENBROCK, Christian
 APPLICANT: BERLIN, Kurt
  TITLE OF INVENTION: Diagnosis of Diseases Associated with
 TITLE OF INVENTION: tumor suppressor genes and oncogenes
 FILE REFERENCE: 5013.1005
  CURRENT APPLICATION NUMBER: US/10/221,714A
  CURRENT FILING DATE: 2003-01-21
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 PRIOR FILING DATE: 2001-03-15
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 PRIOR FILING DATE: 2000-03-15
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Search completed: April 25, 2006, 12:00:54

Job time : 432 secs

SCORE 1.3 BuildDate: 12/06/2005

SCORE Search Results Details for Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rnpbn.

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OM nucleic - nucleic search, using sw model

Run on:

April 25, 2006, 10:54:00; Search time 359 Seconds

(without alignments)

1408.985 Million cell updates/sec

Title:

US-10-088-319-2_COPY_331_455

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Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

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Total number of hits satisfying chosen parameters:

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Minimum DB seg length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Published Applications NA New:*

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4: /SIDS5/ptodata/1/pubpna/PCT NEW PUB.seq:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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	3	30.4	24.3	523	10	US-10-301-480-627931	Sequence 627931,
	4	30.4	24.3	525	9	US-10-301-480-14520	Sequence 14520, A
	. 5	30.4	24.3	525	9	US-10-301-480-14521	Sequence 14520, A
	6	30.4	24.3	525	10	US-10-301-480-627929	Sequence 627929,
	7	30.4	24.3	525	10	US-10-301-480-627930	-
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	9.	30	24.0	525	9	US-10-301-480-14519	Sequence 14519, A
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ALIGNMENTS

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; Publication No. US20060019272A1
; GENERAL INFORMATION:
  APPLICANT: Geraci, Mark
  APPLICANT: Bull, Todd
  APPLICANT: Voelkel, Norbert
  APPLICANT: Coldren, Chris
  TITLE OF INVENTION: Diagnosis of Disease and Monitoring of Therapy Using Gene
  TITLE OF INVENTION: Expression Analysis of Peripheral Blood Cells
  FILE REFERENCE: 2848-54
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  CURRENT FILING DATE: 2005-05-03
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  PRIOR FILING DATE: 2004-05-03
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   ORGANISM: Homo sapiens
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; Publication No. US20060057564A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
  TITLE OF INVENTION: in the Human Genome
  FILE REFERENCE: 108827.137
  CURRENT APPLICATION NUMBER: US/10/301,480
  CURRENT FILING DATE: 2002-11-21
  PRIOR APPLICATION NUMBER: US 10/215,598
  PRIOR FILING DATE: 2002-08-09
  PRIOR APPLICATION NUMBER: US 60/311,695
  PRIOR FILING DATE: 2001-08-10
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; Publication No. US20060057564A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
  TITLE OF INVENTION: in the Human Genome
  FILE REFERENCE: 108827.137
  CURRENT APPLICATION NUMBER: US/10/301,480
  CURRENT FILING DATE: 2002-11-21
  PRIOR APPLICATION NUMBER: US 10/215,598
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; PRIOR APPLICATION NUMBER: US 60/311,695
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; Publication No. US20060057564A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
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TITLE OF INVENTION: in the Human Genome
  FILE REFERENCE: 108827.137
  CURRENT APPLICATION NUMBER: US/10/301,480
  CURRENT FILING DATE: 2002-11-21
  PRIOR APPLICATION NUMBER: US 10/215,598
 PRIOR FILING DATE: 2002-08-09
 PRIOR APPLICATION NUMBER: US 60/311,695
; PRIOR FILING DATE: 2001-08-10
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; Publication No. US20060057564A1
; GENERAL INFORMATION:
 APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
  TITLE OF INVENTION: in the Human Genome
  FILE REFERENCE: 108827.137
  CURRENT APPLICATION NUMBER: US/10/301,480
  CURRENT FILING DATE: 2002-11-21
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  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
  TITLE OF INVENTION: in the Human Genome
  FILE REFERENCE: 108827.137
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  APPLICANT: Wang, David G.
 TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
 TITLE OF INVENTION: in the Human Genome
 FILE REFERENCE: 108827.137
 CURRENT APPLICATION NUMBER: US/10/301,480
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Qу
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RESULT 8
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; Publication No. US20040181048A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single
  TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.135
  CURRENT APPLICATION NUMBER: US/09/925,065A
  CURRENT FILING DATE: 2001-08-08
  PRIOR APPLICATION NUMBER: US 60/243,096
  PRIOR FILING DATE: 2000-10-24
  PRIOR APPLICATION NUMBER: US 60/252,147
  PRIOR FILING DATE: 2000-11-20
  PRIOR APPLICATION NUMBER: US 60/250,092
  PRIOR FILING DATE: 2000-11-30
  PRIOR APPLICATION NUMBER: US 60/261,766
  PRIOR FILING DATE: 2001-01-16
  PRIOR APPLICATION NUMBER: US 60/289,846
  PRIOR FILING DATE: 2001-05-09
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GENERAL INFORMATION:
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  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
  TITLE OF INVENTION: in the Human Genome
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; Publication No. US20060057564A1
; GENERAL INFORMATION:
 APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
 TITLE OF INVENTION: in the Human Genome
 FILE REFERENCE: 108827.137
  CURRENT APPLICATION NUMBER: US/10/301,480
  CURRENT FILING DATE: 2002-11-21
  PRIOR APPLICATION NUMBER: US 10/215,598
  PRIOR FILING DATE: 2002-08-09
  PRIOR APPLICATION NUMBER: US 60/311,695
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                              1; Mismatches
                                              46:
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Qу
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGAACTACGAAGCACATTTT 61
```

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Db
         283 AAAAAAAAAAAAAGATTAGGTAAAATGACTGTTGAAAGGAATGAGTAAGGCACACATTTC 342
Qy
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCC 105
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                     1 11
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RESULT 11
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; Sequence 467966, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
   TITLE OF INVENTION: in the Human Genome
  FILE REFERENCE: 108827.137
  CURRENT APPLICATION NUMBER: US/10/301,480
  CURRENT FILING DATE: 2002-11-21
  PRIOR APPLICATION NUMBER: US 10/215,598
  PRIOR FILING DATE: 2002-08-09
  PRIOR APPLICATION NUMBER: US 60/311,695
  PRIOR FILING DATE: 2001-08-10
 NUMBER OF SEQ ID NOS: 1226818
  SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 467966
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   ORGANISM: Homo sapien
US-10-301-480-467966
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  Best Local Similarity 64.7%; Pred. No. 6;
 Matches 44; Conservative
                              0; Mismatches
                                             24;
                                                   Indels
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         331 AAATTAAGAACGATAACTTAAGGTATTTGGTGGAAGAAATTTCTAAGCAGCAAAGCATTC 272
Db
          64 AGGAAGTG 71
Οv
             1 11 111
        271 AAGATGTG 264
RESULT 12
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; Sequence 1081375, Application US/10301480
; Publication No. US20060057564A1
; GENERAL INFORMATION:
; APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identifiction and Mapping of Single Nucleotide Polymorphisms
  TITLE OF INVENTION: in the Human Genome
  FILE REFERENCE: 108827.137
  CURRENT APPLICATION NUMBER: US/10/301,480
  CURRENT FILING DATE: 2002-11-21
  PRIOR APPLICATION NUMBER: US 10/215,598
  PRIOR FILING DATE: 2002-08-09
  PRIOR APPLICATION NUMBER: US 60/311,695
  PRIOR FILING DATE: 2001-08-10
  NUMBER OF SEQ ID NOS: 1226818
; SOFTWARE: FastSEQ for Windows Version 4.0
; SEQ ID NO 1081375
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LENGTH: 577
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         44; Conservative
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                 Db
         334 AAATTAAGAACGATAACTTAAGGTATTTGGTGGAAGAAATTTCTAAGCAGCAAAGCATTC 275
          64 AGGAAGTG 71
Qу
             1 11 111
         274 AAGATGTG 267
Db
RESULT 13
US-09-925-065A-400790/c
; Sequence 400790, Application US/09925065A
 Publication No. US20040181048A1
 GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single
  TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.135
  CURRENT APPLICATION NUMBER: US/09/925,065A
  CURRENT FILING DATE: 2001-08-08
  PRIOR APPLICATION NUMBER: US 60/243,096
  PRIOR FILING DATE: 2000-10-24
  PRIOR APPLICATION NUMBER: US 60/252,147
  PRIOR FILING DATE: 2000-11-20
  PRIOR APPLICATION NUMBER: US 60/250,092
  PRIOR FILING DATE: 2000-11-30
  PRIOR APPLICATION NUMBER: US 60/261,766
  PRIOR FILING DATE: 2001-01-16
  PRIOR APPLICATION NUMBER: US 60/289,846
  PRIOR FILING DATE: 2001-05-09
  NUMBER OF SEQ ID NOS: 957086
  SOFTWARE: FastSEQ for Windows Version 4.0
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Db
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Db
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RESULT 14

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US-10-857-780-1
; Sequence 1, Application US/10857780
; Publication No. US20050272043A1
; GENERAL INFORMATION:
  APPLICANT: ROTH, RICHARD B.
  APPLICANT: BRAUN, ANDREAS
  APPLICANT: KAMMERER, STEFAN M.
  APPLICANT: NELSON, MATTHEW ROBERTS
  APPLICANT: RENELAND, RIKARD HENRY
  APPLICANT: HOYAL-WRIGHTSON, CAROLYN R.
  TITLE OF INVENTION: METHODS FOR IDENTIFYING RISK OF BREAST CANCER AND TREATMENTS
  TITLE OF INVENTION: THEREOF
  FILE REFERENCE: SEQ-4069-CP
  CURRENT APPLICATION NUMBER: US/10/857,780
  CURRENT FILING DATE: 2004-05-28
  PRIOR APPLICATION NUMBER: 10/723,681
 PRIOR FILING DATE: 2003-11-25
 PRIOR APPLICATION NUMBER: 60/490,234
 PRIOR FILING DATE: 2003-07-24
  PRIOR APPLICATION NUMBER: 60/525,239
  PRIOR FILING DATE: 2003-11-25
  NUMBER OF SEQ ID NOS: 4962
  SOFTWARE: PatentIn version 3.2
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   ORGANISM: Homo sapiens
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   NAME/KEY: misc_feature
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   LOCATION: (24675)..(24676)
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   FEATURE:
   NAME/KEY: misc_feature
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US-10-857-780-1
 Query Match
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                                                   Length 92600;
 Best Local Similarity
                        56.2%; Pred. No. 61;
 Matches
          54; Conservative
                               0; Mismatches
                                               42;
                                                    Indels
                                                              0;
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Qу
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             Db
        3143 CCAAAACAAAGAAACTAAAAATAAGCATTCGGATTTGTTAGGGGGGCGACAAGGGAGGCAC 3202
          61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTT 96
Qу
             Db
        3203 TCCAGGATCTGTGGACTCCCCACTTTGTTCTGTCCT 3238
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RESULT 15
US-09-925-065A-638583
; Sequence 638583, Application US/09925065A
; Publication No. US20040181048A1
; GENERAL INFORMATION:
  APPLICANT: Wang, David G.
  TITLE OF INVENTION: Identification and Mapping of Single
  TITLE OF INVENTION: Nucleotide Polymorphisms in the Human Genome
  FILE REFERENCE: 108827.135
  CURRENT APPLICATION NUMBER: US/09/925,065A
  CURRENT FILING DATE: 2001-08-08
  PRIOR APPLICATION NUMBER: US 60/243,096
  PRIOR FILING DATE: 2000-10-24
  PRIOR APPLICATION NUMBER: US 60/252,147
  PRIOR FILING DATE: 2000-11-20
  PRIOR APPLICATION NUMBER: US 60/250,092
  PRIOR FILING DATE: 2000-11-30
  PRIOR APPLICATION NUMBER: US 60/261,766
  PRIOR FILING DATE: 2001-01-16
  PRIOR APPLICATION NUMBER: US 60/289,846
  PRIOR FILING DATE: 2001-05-09
  NUMBER OF SEQ ID NOS: 957086
  SOFTWARE: FastSEQ for Windows Version 4.0
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   ORGANISM: Homo sapiens
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 Matches
          61; Conservative
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Qу
             Db
          84 CCAAAGATATACAAGTAACCAGTAAGCACAGAAAAAAAACAGCATAACTAATTATTAAAG 143
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Qу
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Search completed: April 25, 2006, 12:06:58
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Job time: 359 secs

SCORE 1.3 BuildDate: 12/06/2005

SCORE Search Results Details for Application 10088319 and Search Result us-10-088-319-2_copy_331_455.rst.

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GenCore version 5.1.7 Copyright (c) 1993 - 2006 Biocceleration Ltd.

OM nucleic - nucleic search, using sw model

Run on:

April 25, 2006, 10:08:11; Search time 1987 Seconds

(without alignments)

2943.324 Million cell updates/sec

Title:

US-10-088-319-2 COPY 331 455

Perfect score:

Sequence:

1 caaaaacaaaaacctttac.....tgagtaaggtggccactttg 125

Scoring table: IDENTITY NUC

Gapop 10.0 , Gapext 1.0

Searched:

41078325 seqs, 23393541228 residues

Total number of hits satisfying chosen parameters:

82156650

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

EST:*

1: qb est1:* 2: gb est2:*

3: gb_est3:*

4: qb htc:*

5: gb_est4:*

6: qb est5:*

7: gb_est6:*

8: gb est7:*

9: gb gss1:*

10: gb gss2:*

11: gb_gss3:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

			용				
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	2	105.4	84.3	665	8	DN995162	DN995162 TC112886
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С	4	31.8	25.4	578	2	BI291532	BI291532 UI-R-DM0-
	5	31.6	25.3	480	10	CE391194	CE391194 tigr-gss-
С	6	31.4	25.1.	788	9	BZ459393	BZ459393 BONIE51TF
С	7	31.4	25.1	1032	11	CNS05L3M	AL342283 Tetraodon
С	8	31.2	25.0	529	2	BI291766	BI291766 UI-R-DM0-
	9	31.2	25.0	579	7	CF918988	CF918988 Bflor531.
С	10	31.2	25.0	627	2	BI285373	BI285373 UI-R-DBO-
С	11	31.2	25.0	630	2	BI279004	BI279004 UI-R-DB0-
С	12	31.2	25.0	655	2	BI285082	BI285082 UI-R-DB0-
	13	31	24.8	669	10	CW685817	. CW685817 OG BBa004
С	14	30.8	24.6	422	3	BI527837	BI527837 1024084H1
С	15	30.8	24.6	452	3	BI999848	BI999848 1031079E0
С	16	30.8	24.6	508	3	BI995666	BI995666 1031029D0
С	17	30.8	24.6	553	5	BU652076	BU652076 1112097D0
С	18	30.8	24.6	601	3	BM003194	`BM003194 1031109A1
C	19	30.8	24.6	613	3	BI723641	BI723641 1031067D0
С	20	30.8	24.6	638	3	BI723640 .	BI723640 1031067D0
С	21	30.8	24.6	639	3	BI724908	BI724908 1031075H0
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С	26	30.6	24.5	432	9	BZ768094	BZ768094 SALK 1397
С	27	30.6	24.5	474	6	CB407550	CB407550 IPG030C06
	28	30.6	24.5	478	7	CF962820	CF962820 8997rsice
	29	30.6	24.5	634	5	BU437831	BU437831 604145016
С	30	30.6	24.5	1017	3	BQ050581	BQ050581 AGENCOURT
	31	30.6	24.5	1550	10	CL019408	CL019408 CH216-5D1
	32	30.4	24.3	495	7	CN679199	CN679199 E0121C11-
	33	30.4	24.3	552	7	CN681667	CN681667 E0153F05-
С	34	30.4	24.3	594	3	BJ017277	BJ017277 BJ017277
С	35	30.4	24.3	741	8	DR103237	DR103237 JHU027E09
С	36	30.4	24.3	866	7	CN172789	CN172789 AGENCOURT
С	37	30.2	24.2	433	9	AZ618267	AZ618267 1M0449I20
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	41	30.2	24.2	864	6	CD379455	CD379455 PTMM04624
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ALIGNMENTS

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LOCUS
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ACCESSION CK834247
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VERSION
           CK834247.1 GI:45064536
KEYWORDS
           EST.
           Bos taurus (cow)
SOURCE
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           Mammalia; Eutheria; Laurasiatheria; Cetartiodactyla; Ruminantia;
           Pecora; Bovidae; Bovinae; Bos.
              (bases 1 to 740)
REFERENCE
           Baumann, R.G., Baldwin, R.L., Sonstegard, T.S., Van Tassell, C.P. and
 AUTHORS
           Matukumalli, L.K.
 TITLE
           Construction and Analysis of a cDNA Library Generated From
           Intestinal Muscle and Epithelial Tissues of Holstein Cattle
           Unpublished (2004)
 JOURNAL
           Contact: Richard G. Baumann
COMMENT
           Bovine Functional Genomics Lab
           ANRI
           BLDG 162: BARC-EAST, Beltsville, MD 20705, USA
           Tel: 3015048604
           Fax: 3015048744
           Email: rbaumann@anri.barc.usda.gov
           Single pass sequencing. Bases called and trimmed with phred
           0.000925 using options -trim alt '' -trim fasta. Vector identified
           by cross match using options -minmatch 12 -minscore 12
           Plate: 2 row: M column: 16
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           High quality sequence stop: 740.
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                   NotI; Site 2: EcoRI; Normalized cow cDNA intestinal
                   library in pCMVsport6.1, constructed from equimolar mRNA
                   pools derived from 5 sources, 4 lactating intestinal, 1
                   neonatal intestinal 4/5 Lactating, Proximal Duodenum,
                   Jejunum, Distal Ileum, Colon, 1/5 Neonatal, Proximal
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                               Score 119.2; DB 7;
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Qу
             Db
         162 CCAGGAAGTGTGGGCTGCGACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 221
         122 TTTG 125
Qy
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 $I \cup I \cup$

Query Match

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LOCUS
            DN995162
                                      665 bp
                                                mRNA
                                                        linear
                                                                 EST 17-MAY-2005
            TC112886 Human adult whole brain, large insert, pCMV expression
DEFINITION
            library Homo sapiens cDNA clone TC112886 5' similar to Homo sapiens
            CD40 ligand (TNF superfamily, member 5, hyper-IgM syndrome)
            (CD40LG), mRNA sequence.
ACCESSION
            DN995162
VERSION
            DN995162.1 GI:66254993
KEYWORDS
            EST.
SOURCE
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  ORGANISM Homo sapiens
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Primates; Catarrhini;
            Hominidae; Homo.
REFERENCE
            1 (bases 1 to 665)
  AUTHORS
            Birkett, C., Cho, J., Gau, Y., Hamer, R., Kelly, S., Kovacs, K., Liu, L.,
            Liu, X., Porter, J., Sachs, A., Shu, Y., Sun, Z., Wong, J., Wu, M.,
            Zhang, X., Jay, G. and He, W.
  TITLE
            High-throughput cloning of full-length human cDNAs directly from
            cDNA libraries optimized for large and rare transcripts
  JOURNAL
            Unpublished (2005)
            Contact: Kovacs, KF
COMMENT
            High Throughput cDNA Cloning
            OriGene Technologies, Inc. ( www.origene.com )
            6 Taft Court, Suite 100, Rockville, MD 20850, USA
            Tel: 301 340 3188
            Fax: 301 340 8606
            Email: cDNA@origene.com
            This EST submission is part of an on-going human full-length
            cloning project at OriGene Technologies, Inc.
            Please contact OriGene for access.
            OriGene Technologies, Inc.
            6 Taft Ct. Suite 100
            Rockville, MD 20850
            Tel: (301) 340-3188
            http://www.origene.com
            Seq primer: pCMV6 5prime forward vector primer, OriGene
            Technologies Inc.
FEATURES
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                     expression library"
                     /note="Organ: Brain; Vector: pCMV6-XL5; Site 1: EcoR1;
                     Site 2: Xhol/Sall compatible end ligatio; Oligo-dT primed
                     reverse transcription optimized for large and GC rich mRNA
                     transcripts, cDNA size selection, optimized ligation for
                     large inserts into mammalian expression vector, random
                     clones selected for end sequence verification of
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ORIGIN
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Qy
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Db
          79 CAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCACTTTG 125
Qy
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Db
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                                                             EST 04-NOV-2002
           CA339153
                                                     linear
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           IMAGE: 5621810 5', mRNA sequence.
ACCESSION
           CA339153
VERSION
           CA339153.1 GI:24557251
KEYWORDS
           EST.
SOURCE
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 ORGANISM Rattus norvegicus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
           Sciurognathi; Muroidea; Muridae; Murinae; Rattus.
REFERENCE
           1 (bases 1 to 650)
 AUTHORS
           NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE
           National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
           Tumor Gene Index
 JOURNAL
           Unpublished (1997)
COMMENT
           Contact: Robert Strausberg, Ph.D.
           Email: cgapbs-r@mail.nih.gov
           cDNA Library Preparation:
            cDNA Library Arrayed by: The I.M.A.G.E. Consortium/LLNL
            DNA Sequencing by: National Institutes of Health Intramural
           Sequencing Center (NISC)
            Clone distribution: NCI-CGAP clone distribution information can be
           found through the I.M.A.G.E. Consortium/LLNL at:
           info@image.llnl.gov
           Plate: LLAM12446 row: H column: 3
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                    NotI; Site 2: EcoRV; Cloned unidirectionally. Primer:
                    Oligo dT. Pool of 2 primary libraries: NCI CGAP Pr46
                    (ventral prostate from 10 wk male, average insert size 2
                    kb) and NCI_CGAP_Pr47 (dorsolateral prostate from 10 wk
                    male, average insert size 2 kb). Constructed by
                    Invitrogen. Note: this is a NCI CGAP Library."
ORIGIN
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                                 Pred. No. 8.1;
           67; Conservative 0; Mismatches
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                                                                           0;
           Qу
                        1 11 111 11 11111
                                                            1111
                                                                   11
Db
         209 AAACAACAGTGACTTGTATCTAAGTCGTGTGGTGGAAGTGAAGACAGTGAAGACAATCCA 268
          62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qу
                         -11
                                           1 1 1 1 111 1 11 11
Db
         269 ACAGCAGATAGAGGCTGGAAAGACCTTGTTATTTGATGTGATCTTAAGCAAAACAGCATG 328
         122 TTTG 125
Οv
             1111
Db
         329 TTTG 332
RESULT 4
BI291532/c
LOCUS
           BI291532
                                    578 bp
                                             mRNA
                                                     linear
                                                              EST 19-JUL-2001
DEFINITION UI-R-DMO-cis-n-19-0-UI.sl UI-R-DMO Rattus norvegicus cDNA clone
           UI-R-DMO-cis-n-19-0-UI 3', mRNA sequence.
           BI291532
ACCESSION
           BI291532.1 GI:14951161
VERSION
KEYWORDS
           EST.
SOURCE
           Rattus norvegicus (Norway rat)
 ORGANISM Rattus norvegicus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
           Sciurognathi; Muroidea; Muridae; Murinae; Rattus.
REFERENCE
              (bases 1 to 578)
 AUTHORS
           Bonaldo, M.F., Lennon, G. and Soares, M.B.
 TITLE
           Normalization and subtraction: two approaches to facilitate gene
           discovery
           Genome Res. 6 (9), 791-806 (1996)
 JOURNAL 
  PUBMED
           8889548
COMMENT
           Contact: Soares, MB
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           The sequence contained an oligo-dT track that was present in the
           oligonucleotide that was used to prime the synthesis of first
           strand cDNA and therefore this may represent a bonafide poly A
           tail. The sequence tag present in the cDNA between the NotI site
           and the oligo-dT track served to verify it as a clone from the
           non-normalized rat prostate library cDNA Library Preparation: M.B.
          Soares Lab Clone distribution: clones will be available through
           Research Genetics (www.resgen.com) The following repetitive
           elements were found in this cDNA sequence: 1-21,
           >AT_rich#Low_complexity
           Seq primer: M13 Forward
           POLYA=Yes.
FEATURES
                    Location/Qualifiers
    source
                    1. .578
                    /organism="Rattus norvegicus"
                    /mol type="mRNA"
                    /strain="Sprague-Dawley"
                    /db_xref="taxon:10116"
```

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/clone="UI-R-DMO-cis-n-19-0-UI"
                     /dev stage="ADULT"
                     /lab_host="DH10B (Life Technologies)"
                     /clone lib="UI-R-DM0"
                     /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                     polylinker; Site_1: Not I; Site_2: Eco RI; The UI-R-DMO
                     library is a non-normalized Rat Prostate library
                     constructed in pT3T7 PAC vector according to the procedure
                     described by Bonaldo, Lennon & Soares (Genome Research
                     Genome 6: 791-806, 1996). The oligonucleotide used to
                     prime first strand synthesis contained the sequence tag
                     CCAGG between the Not I cloning site and dT18 stretch.
                     TAG TISSUE=rat prostate
                     TAG LIB=UI-R-DMO
                     TAG SEQ=CCAGG"
                          25.4%; Score 31.8; DB 2; Length 578;
 Best Local Similarity
                         53.2%;
                                 Pred. No. 17;
                                0; Mismatches
           66; Conservative
                                                  58; Indels
                                                                             0:
                                                                     Gaps
           2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
                         1 11 111 11 11111
          537 AAACAACAGTGACTTGTATNTAAGTCGTGTGGTGGAAGTGAAGACAGTGAAGACAATCCA 478
           62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
                                             1 1 1 1 11 1 1 11
                         11
          477 ACAGCAGATAGAGGCTGGAAAGACCTTGTTATTTGATGTGATCTTAGGCAAAACAGCATG 418
         122 TTTG 125
          417 TTTG 414
           CE391194
                                     480 bp
                                               DNA
                                                       linear
                                                                GSS 27-SEP-2003
DEFINITION tigr-gss-dog-17000334460145 Dog Library Canis familiaris genomic,
           genomic survey sequence.
           CE391194
           CE391194.1 GI:36630810
           Canis familiaris (dog)
 ORGANISM Canis familiaris
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Laurasiatheria; Carnivora; Fissipedia; Canidae;
               (bases 1 to 480)
           Kirkness, E.F., Bafna, V., Halpern, A.L., Levy, S., Remington, K.,
           Rusch, D.B., Delcher, A.L., Pop, M., Wang, W., Fraser, C.M. and
           The dog genome: survey sequencing and comparative analysis
           Science 301 (5641), 1898-1903 (2003)
           14512627
           Contact: Kirkness EF
            The Institute for Genomic Research
           Department of Eukaryotic Genomics, TIGR, 9712 Medical Center Drive,
           Rockville, MD 20850, USA
            Tel: 301-838-0200
            Fax: 301-838-0208
           Email: ekirknes@tigr.org
```

ORIGIN

Qy

Db

Qy

Db

Qy

Db

RESULT 5 CE391194 LOCUS

ACCESSION

VERSION

SOURCE

KEYWORDS

REFERENCE

TITLE

COMMENT

JOURNAL

PUBMED

AUTHORS

Query Match

GSS.

Canis.

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Class: shotqun.
                     Location/Qualifiers
FEATURES
                     1. .480
     source
                     /organism="Canis familiaris"
                     /mol type="genomic DNA"
                     /strain="Standard Poodle"
                     /db xref="taxon:9615"
                     /clone_lib="Dog Library"
                     /note="Site_1: BstXI; Libraries were prepared from
                     peripheral blood"
ORIGIN
                                  Score 31.6; DB 10;
  Query Match
                          25.3%;
                                                        Length 480;
  Best Local Similarity
                          55.5%;
                                  Pred. No. 19;
  Matches
            61; Conservative
                                     Mismatches
                                 0;
                                                   49;
                                                        Indels
                                                                              0:
                                                                      Gaps
Qy
            2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
              111 1 1 1 11 1111
                                  7 AAATAAATAGAATCTTTAAAAAAATTTTTTTTTTTCAAGTGAAAACTTTGGAAAACTTTTA 66
Db
           62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTA 111
Qу
                                               11111 | 1 | 11111 |
                         | | | |
              11111
                               - 1
                                  1 11
           67 CCTGACATCAGGGGCATGACAGGTTCAAAATACTTAAAGACTTCTGAGAA 116
Db
RESULT 6
BZ459393/c
LOCUS
            BZ459393
                                     788 bp
                                               DNA
                                                                 GSS 13-DEC-2002
                                                        linear
            BONIE51TF BO 1.6 2 KB tot Brassica oleracea genomic clone BONIE51,
DEFINITION
            genomic survey sequence.
ACCESSION
            BZ459393
VERSION
            BZ459393.1 GI:26740861
KEYWORDS
            GSS.
SOURCE
            Brassica oleracea
  ORGANISM Brassica oleracea
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; eudicotyledons; core eudicotyledons;
            rosids; eurosids II; Brassicales; Brassicaceae; Brassica.
REFERENCE
               (bases 1 to 788)
  AUTHORS
            Ayele, M., Haas, B.J., Kumar, N., Wu, H., Xiao, Y., Van Aken, S.,
            Utterback, T.R., Wortman, J.R., White, O.R. and Town, C.D.
  TITLE
            Whole genome shotgun sequencing of Brassica oleracea and its
            application to gene discovery and annotation in Arabidopsis
            Genome Res. 15 (4), 487-495 (2005)
  JOURNAL
            15805490
   PUBMED
COMMENT
            Other GSSs: BONIE51TR
            Contact: Chris Town
            TTGR
            9712 Medical Center Drive, Rockville, MD 20850, USA.
            Tel: 301-838-3523
            Fax: 301-838-0208
            Email: cdtown@tigr.org
            DNA is from a doubled haploid provided by Tom Osborn.
            Seg primer: TF
            Class: sheared ends.
                     Location/Qualifiers
FEATURES
                     1. .788
     source
                     /organism="Brassica oleracea"
                     /mol type="genomic DNA"
                     /strain="TO1000DH3"
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/db_xref="taxon:3712"

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/clone="BONIE51"
                     /clone lib="BO 1.6 2 KB tot"
                     /note="Vector: pHOS1; Site_1: BstXI; 1.6-2 kb sheared
                     total DNA inserted into pHOS1 using BstXI linkers"
ORIGIN
 Query Match
                          25.1%;
                                  Score 31.4; DB 9;
                                                      Length 788;
  Best Local Similarity
                          61.7%;
                                  Pred. No. 25;
            50; Conservative
                                 0; Mismatches
                                                   31;
                                                        Indels
                                                                              0:
            1 CAAAAACAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTT 60
Qy
                   11 11
                                                  11
Db
          710 CAAACGCATAAATTCTTTATGTGGAGTATTCCTATGGAGAGGTAACATTGAAGAACATCA 651
Qу
           61 TCCAGGAAGTGTGGGCTGCAA 81
              1 111 11 111 1 11 11
Db
          650 TACAGCGAGGGTGAGTTGGAA 630
RESULT 7
CNS05L3M/c
LOCUS
            CNS05L3M
                                    1032 bp
                                               DNA
                                                        linear
                                                                 GSS 01-SEP-2000
DEFINITION Tetraodon nigroviridis genome survey sequence T7 end of clone
            051F03 of library A from Tetraodon nigroviridis, genomic survey
            sequence.
ACCESSION
            AL342283
VERSION
            AL342283.1 GI:8236041
KEYWORDS
            GSS; genome survey sequence.
SOURCE
            Tetraodon nigroviridis
  ORGANISM
           Tetraodon nigroviridis
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Actinopterygii; Neopterygii; Teleostei; Euteleostei; Neoteleostei;
            Acanthomorpha; Acanthopterygii; Percomorpha; Tetraodontiformes;
            Tetradontoidea; Tetraodontidae; Tetraodon.
REFERENCE
            1
  AUTHORS
            Roest Crollius, H., Jaillon, O., Dasilva, C., Bouneau, L., Fisher, C.,
            Bernot, A., Fizames, C., Wincker, P., Brottier, P., Quetier, F.,
            Saurin, W. and Weissenbach, J.
  TITLE
            Estimate of human gene number provided by genome-wide analysis
            using Tetraodon nigroviridis DNA sequence
  JOURNAL
            Nat. Genet. 25 (2), 235-238 (2000)
   PUBMED
            10835645
REFERENCE
 AUTHORS
            Roest Crollius, H., Jaillon, O., Dasilva, C., Ozouf-Costaz, C.,
            Fizames, C., Fischer, C., Bouneau, L., Billault, A., Quetier, F.,
            Saurin, W., Bernot, A. and Weissenbach, J.
  TITLE
            Characterization and repeat analysis of the compact genome of the
            freshwater pufferfish Tetraodon nigroviridis
  JOURNAL
            Genome Res. 10 (7), 939-949 (2000)
   PUBMED
            10899143
            3 (bases 1 to 1032)
REFERENCE
 AUTHORS
           Genoscope.
  TITLE
           Direct Submission
  JOURNAL
            Submitted (12-APR-2000) Genoscope - Centre National de Sequencage :
            BP 191 91006 EVRY cedex - FRANCE (E-mail : seqref@genoscope.cns.fr
            Web : www.genoscope.cns.fr)
COMMENT
            This sequence is a single read and was generated as part of a large
            scale clone-end sequencing project of the Tetraodon nigroviridis
            genome. For more information, please take a look at
            http://www.genoscope.cns.fr/Tetraodon.
                     Location/Qualifiers
FEATURES
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1. .1032
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                     /organism="Tetraodon nigroviridis"
                     /mol type="genomic DNA"
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                     /clone="051F03"
                     /clone lib="A"
                     /note="Genoscope sequence ID : COAAO51CC02C1
                     end : T7"
ORIGIN
 Query Match
                                 Score 31.4; DB 11;
                         25.1%;
                                                       Length 1032;
 Best Local Similarity
                         51.0%; Pred. No. 27;
           53; Conservative
                                0; Mismatches
                                                  51;
                                                       Indels
                                                                    Gaps
            4 AAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTTCC 63
Qy
              11 1111 11 11 1
                                    -11 1 1
                                                            Db
          619 AAGGAAAAANCCCNTATGAAGTCNTTTTTCANGNNNGNAAAAANANNCANNNCATTTTCT 560
          64 AGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTG 107
Qy
                4 411 1 11 11 111
                                         Db
          559 CTGTTGTGAGAGCCTAAAAAAATGTATCATGTTTAAAATTNCTG 516
RESULT 8
BI291766/c
           BI291766
LOCUS
                                    529 bp
                                              mRNA
                                                       linear
                                                               EST 19-JUL-2001
DEFINITION UI-R-DMO-cis-n-18-0-UI.sl UI-R-DMO Rattus norvegicus cDNA clone
           UI-R-DMO-cis-n-18-0-UI 3', mRNA sequence.
           BI291766
ACCESSION
           BI291766.1 GI:14951628
VERSION
KEYWORDS
           EST.
SOURCE
           Rattus norvegicus (Norway rat)
  ORGANISM Rattus norvegicus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
           Sciurognathi; Muroidea; Muridae; Murinae; Rattus.
REFERENCE
               (bases 1 to 529)
 AUTHORS
           Bonaldo, M.F., Lennon, G. and Soares, M.B.
 TITLE
           Normalization and subtraction: two approaches to facilitate gene
           discovery
  JOURNAL
           Genome Res. 6 (9), 791-806 (1996)
  PUBMED
           8889548
COMMENT
           Contact: Soares, MB
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           The sequence contained an oligo-dT track that was present in the
           oligonucleotide that was used to prime the synthesis of first
           strand cDNA and therefore this may represent a bonafide poly A
           tail. The sequence tag present in the cDNA between the NotI site
            and the oligo-dT track served to verify it as a clone from the
           non-normalized rat prostate library cDNA Library Preparation: M.B.
           Soares Lab Clone distribution: clones will be available through
           Research Genetics (www.resgen.com)
           Seq primer: M13 Forward
            POLYA=Yes.
FEATURES
                    Location/Qualifiers
     source
                    1. .529
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/organism="Rattus norvegicus"
                     /mol type="mRNA"
                     /strain="Sprague-Dawley"
                     /db_xref="taxon:10116"
                     /clone="UI-R-DMO-cis-n-18-0-UI"
                     /dev_stage="ADULT"
                     /Tab host="DH10B (Life Technologies)"
                     /clone lib="UI-R-DM0"
                     /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                    polylinker; Site 1: Not I; Site 2: Eco RI; The UI-R-DMO
                     library is a non-normalized Rat Prostate library
                     constructed in pT3T7 PAC vector according to the procedure
                    described by Bonaldo, Lennon & Soares (Genome Research
                    Genome 6: 791-806, 1996). The oligonucleotide used to
                    prime first strand synthesis contained the sequence tag
                    CCAGG between the Not I cloning site and dT18 stretch.
                     TAG TISSUE=rat prostate
                     TAG_LIB=UI-R-DMO
                     TAG SEQ=CCAGG"
ORIGIN
                         25.0%;
  Query Match .
                                 Score 31.2; DB 2;
                                                     Length 529;
  Best Local Similarity
                         53.2%;
                                 Pred. No. 26;
           66; Conservative
                                0; Mismatches
                                                      Indels
Qу
            2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
                         1 11 111 11 11111
                                                              1111
Db
          527 AAACAACAGTGACTTGTATCTAAGTCGTGTGGTGGAAGTGAAGACAGTGAAGACAATCCA 468
Qу
           62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
                          - 11
                                            11
Db
          467 ACAGCAGATAGAGGCTGGAAAGACCTTGTTATTTGATGTGATCTTAGGCAAAACAGCATG 408
          122 TTTG 125
Qy
              ++++
          407 TTTG 404
Db
RESULT 9
CF918988
LOCUS
           CF918988
                                    579 bp
                                              mRNA
                                                      linear
                                                               EST 05-NOV-2003
DEFINITION Bflor531.000073 Amphioxus 26 hrs cDNA library (Name convention:
           BFL26 or MPMGp531) Branchiostoma floridae cDNA clone
           MPMGp531J1510;BFL26_10J15 5', mRNA sequence.
           CF918988
ACCESSION
VERSION
           CF918988.1 GI:38190190
KEYWORDS
SOURCE
           Branchiostoma floridae (Florida lancelet)
  ORGANISM Branchiostoma floridae
           Eukaryota; Metazoa; Chordata; Cephalochordata; Branchiostomidae;
           Branchiostoma.
REFERENCE
           1 (bases 1 to 579)
 AUTHORS
           Panopoulou, G., Hennig, S., Groth, D., Krause, A., Poustka, A.J.,
           Herwig, R., Vingron, M. and Lehrach, H.
  TITLE
           New evidence for genome-wide duplications at the origin of
           vertebrates using an amphioxus gene set and completed animal
           genomes
  JOURNAL
           Genome Res. 13 (6A), 1056-1066 (2003)
  PUBMED
           12799346
COMMENT
           Contact: Panopoulou G
           laboratory 145, dept.Lehrach
```

```
Max-Planck-Institut fuer Molekulare Genetik
           Ihnestr. 63-73, D-14195 Berlin, Germany
           Tel: +49 30 8413 1235
           Fax: +49 30 8413 1128
           Email: panopoul@molgen.mpg.de
           The library was characterised by oligonucleotide fingerprinting
           (ONFP) to reduce sequencing redundancy. According to the ONFP
           procedure clones giving the same hybridisation pattern with a
           battery of 200 8mer oligonucleotides are grouped into clusters. One
           clone per cluster is selected for sequencing. The size of each
           cluster is an indicator of the frequency of a transcript in the
           analysed library. The cluster size as well as the coordinates of
           all clones assigned to the same fingerprint cluster as the clone
           from which the above EST is generated is available at the amphioxus
           project site at http://www.molgen.mpg.de/amphioxus.
           Clones and filters are distributed via the Resource Center/Primary
           Database of the German Genome Project (http://www.rzpd.de).
           PCR PRimers
           FORWARD: 5' CCCCAGGCTTTACACTTTATGCTTCCGGCTCG 3' (M13RSP)
           BACKWARD: 5' GCTATTACGCCAGCTGGCGAAAGGGGGGATGTG 3' (M13FSP)
           Insert Length: 1200
                               Std Error: 200.00
           Seq primer: 5'-CCGGTCCGGAATTCCCGGGT-3' psport3/86
           High quality sequence stop: 579.
FEATURES
                    Location/Qualifiers
    source
                    1. .579
                    /organism="Branchiostoma floridae"
                    /mol type="mRNA"
                    /strain="wild type"
                    /db_xref="taxon:7739"
                    /clone="MPMGp531J1510;BFL26 10J15"
                    /tissue_type="whole embryo"
                    /dev stage="26 hrs (neurula stage)"
                    /lab_host="Escherichia coli, XL1 blue"
                    /clone lib="Amphioxus 26 hrs cDNA library (Name
                    convention: BFL26 or MPMGp531)"
                    /note="Vector: pSport1; Site_1: SalI, KpnI, EcoRI (5');
                    Site_2: NotI, BamHI, HindIII (3'); OligodT primed and
                    directionally cloned in pSport1 vector using a NotI
                    (5'-pGACTAGTTCTAGATCGCGAGCGGCCCC (T)15-3' and a SalI 5'-
                    TCGACCCACGCGTCCG-3'adapters (Gibco BRL)."
ORIGIN
 Query Match
                         25.0%;
                                Score 31.2; DB 7; Length 579;
 Best Local Similarity
                        57.0%;
                                Pred. No. 27;
           57; Conservative
                               0; Mismatches
                                                43;
                                                    Indels
                                                              0; Gaps
                                                                          0;
Qy
           3 AAAACAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTTC 62
             11
                  1 1 11 11 11
                               - 1
                                     11111111
                                              Db
         Qy
          63 CAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAA 102
              Db
         399 AAGCAAGAGAGACAGAAAAGAAGCTGAGCTTTTAACAAA 438
RESULT 10
BI285373/c
LOCUS
           BI285373
                                   627 bp
                                            mRNA
                                                    linear
                                                             EST 19-JUL-2001
DEFINITION UI-R-DB0-byz-c-11-0-UI.sl UI-R-DB0 Rattus norvegicus cDNA clone
           UI-R-DBO-byz-c-11-0-UI 3', mRNA sequence.
ACCESSION
           BI285373
```

```
KEYWORDS
SOURCE
            Rattus norvegicus (Norway rat)
  ORGANISM Rattus norvegicus
            Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
            Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
            Sciurognathi; Muroidea; Muridae; Murinae; Rattus.
REFERENCE
               (bases 1 to 627)
  AUTHORS
            Bonaldo, M.F., Lennon, G. and Soares, M.B.
  TITLE
            Normalization and subtraction: two approaches to facilitate gene
  JOURNAL
            Genome Res. 6 (9), 791-806 (1996)
            8889548
   PUBMED
COMMENT
            Contact: Soares, MB
            Coordinated Laboratory for Computational Genomics
            University of Iowa
            375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
            Tel: 319 335 8250
            Fax: 319 335 9565
            Email: bento-soares@uiowa.edu
            The sequence contained an oligo-dT track that was present in the
            oligonucleotide that was used to prime the synthesis of first
            strand cDNA and therefore this may represent a bonafide poly A
            tail. The sequence tag present in the cDNA between the NotI site
            and the oligo-dT track served to verify it as a clone from the
            non-normalized bladder library cDNA Library Preparation: M.B.
            Soares Lab Clone distribution: clones will be available through
            Research Genetics (www.resgen.com)
            Seq primer: M13 Forward
            POLYA=Yes.
FEATURES
                     Location/Qualifiers
     source
                     1. .627
                     /organism="Rattus norvegicus"
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                     /strain="Sprague-Dawley"
                     /db_xref="taxon:10116"
                     /clone="UI-R-DB0-byz-c-11-0-UI"
                     /dev stage="ADULT"
                     /lab host="DH10B (Life Technologies)"
                     /clone lib="UI-R-DB0"
                     /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                     polylinker; Site_1: Not I; Site_2: Eco RI; The UI-R-DBO
                     library is a non-normalized library constructed from rat
                     bladder tissue. For a detailed description of the library
                     from which this clone was derived, please visit our web
                     site at ratest.eng.uiowa.edu. The subtraction has been
                     previously described in (Bonaldo, Lennon and Soares,
                     Genome Research 6:791-806, 1996)
                     TAG_TISSUE=bladder
                     TAG_LIB=UI-R-DB0
                     TAG SEQ=AGC"
ORIGIN
 Query Match
                          25.0%;
                                 Score 31.2;
                                               DB 2;
                                                      Length 627;
 Best Local Similarity
                          53.2%;
                                 Pred. No. 28;
           66; Conservative
                                 0; Mismatches
                                                  58;
                                                       Indels
                                                                     Gaps
                                                                              0;
Qy
            2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAGAAGACTACGAAGCACATTTT 61
                        1 11 111 11 1111
                                                              1111
          525 AAACAACAGTGACTTGTATCTAAGTCGTGTGGTGGAAGTGAAGACAGTGAAGACAATCCA 466
Db
```

VERSION

.BI285373.1 GI:14938944

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62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qу
                          11
                                             Db
          465 ACAGCAGATAGAGGCTGGAAAGACCTTGTTATTTGATGTGATCTTAGGCAAAACAGCATG 406 .
          122 TTTG 125.
Qy
              1111
          405 TTTG 402
Db
RESULT 11
BI279004/c
LOCUS
           BI279004
                                     630 bp
                                               mRNA
                                                       linear
                                                                EST 19-JUL-2001
DEFINITION UI-R-DB0-byt-h-12-0-UI.sl UI-R-DB0 Rattus norvegicus cDNA clone
           UI-R-DBO-byt-h-12-0-UI 3', mRNA sequence.
ACCESSION
           BI279004
VERSION
           BI279004.1 GI:14926402
KEYWORDS
           EST.
SOURCE
           Rattus norvegicus (Norway rat)
  ORGANISM Rattus norvegicus
           Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
           Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia;
           Sciurognathi; Muroidea; Muridae; Murinae; Rattus.
               (bases 1 to 630)
REFERENCE
 AUTHORS
           Bonaldo, M. F., Lennon, G. and Soares, M.B.
 TITLE
           Normalization and subtraction: two approaches to facilitate gene
           discovery
  JOURNAL
           Genome Res. 6 (9), 791-806 (1996)
           8889548
  PUBMED
COMMENT
           Contact: Soares, MB
           Coordinated Laboratory for Computational Genomics
           University of Iowa
           375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA
           Tel: 319 335 8250
           Fax: 319 335 9565
           Email: bento-soares@uiowa.edu
           The sequence contained an oligo-dT track that was present in the
           oligonucleotide that was used to prime the synthesis of first
           strand cDNA and therefore this may represent a bonafide poly A
           tail. The sequence tag present in the cDNA between the NotI site
           and the oligo-dT track served to verify it as a clone from the
           non-normalized bladder library cDNA Library Preparation: M.B.
           Soares Lab Clone distribution: clones will be available through
           Research Genetics (www.resgen.com)
           Seg primer: M13 Forward
           POLYA=Yes.
FEATURES
                     Location/Qualifiers
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                     /organism="Rattus norvegicus"
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                     /strain="Sprague-Dawley"
                     /db xref="taxon:10116"
                     /clone="UI-R-DB0-byt-h-12-0-UI"
                    /dev stage="ADULT"
                     /lab host="DH10B (Life Technologies)"
                     /clone lib="UI-R-DB0"
                     /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                    polylinker; Site_1: Not I; Site_2: Eco RI; The UI-R-DBO
                    library is a non-normalized library constructed from rat
                    bladder tissue. For a detailed description of the library
                     from which this clone was derived, please visit our web
```

site at ratest.eng.uiowa.edu. The subtraction has been

previously described in (Bonaldo, Lennon and Soares, Genome Research 6:791-806, 1996) TAG TISSUE=bladder TAG LIB=UI-R-DB0 TAG_SEQ=AGC"

Score 31.2; DB 2; Length 630;

ORIGIN

Query Match

Best Local Similarity Pred. No. 28; 53.2%; Matches 66; Conservative 0; Mismatches 58; 0; Indels Gaps 2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61 Qу H + H + H1 11 111 11 11111 - 1 HHHDb 526 AAACAACAGTGACTTGTATCTAAGTCGTGTGGTGGAAGTGAAGACAGTGAAGACAATCCA 467 62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121 Qу -111 | 1 | 1 | 1 | 1 | 1 | 1 Db 466 ACAGCAGATAGAGGCTGGAAAGACCTTGTTATTTGATGTGATCTTAGGCAAAACAGCATG 407 122 TTTG 125 Qу IIIII406 TTTG 403 Db RESULT 12 BI285082/c LOCUS 655 bp BI285082 mRNA linear EST 19-JUL-2001 DEFINITION UI-R-DB0-byw-a-02-0-UI.sl UI-R-DB0 Rattus norvegicus cDNA clone UI-R-DBO-byw-a-02-0-UI 3', mRNA sequence. ACCESSION BI285082 BI285082.1 GI:14938372 VERSION KEYWORDS EST. Rattus norvegicus (Norway rat) SOURCE ORGANISM Rattus norvegicus Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Euarchontoglires; Glires; Rodentia; Sciurognathi; Muroidea; Muridae; Murinae; Rattus. REFERENCE 1 (bases 1 to 655) AUTHORS Bonaldo, M.F., Lennon, G. and Soares, M.B. TITLE Normalization and subtraction: two approaches to facilitate gene discovery JOURNAL Genome Res. 6 (9), 791-806 (1996) PUBMED 8889548 COMMENT Contact: Soares, MB

25.0%;

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375 Newton Road , 4156 MEBRF, Iowa City, IA 52242, USA

Tel: 319 335 8250 Fax: 319 335 9565

Email: bento-soares@uiowa.edu

The sequence contained an oligo-dT track that was present in the oligonucleotide that was used to prime the synthesis of first strand cDNA and therefore this may represent a bonafide poly A tail. The sequence tag present in the cDNA between the NotI site and the oligo-dT track served to verify it as a clone from the non-normalized bladder library cDNA Library Preparation: M.B. Soares Lab Clone distribution: clones will be available through Research Genetics (www.resgen.com) The following repetitive elements were found in this cDNA sequence: 1-21,

>AT_rich#Low_complexity Seq primer: M13 Forward POLYA=Yes.

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                    /dev stage="ADULT"
                    /lab host="DH10B (Life Technologies)"
                    /clone lib="UI-R-DB0"
                    /note="Vector: pT7T3D-Pac (Pharmacia) with a modified
                    polylinker; Site 1: Not I; Site 2: Eco RI; The UI-R-DBO
                    library is a non-normalized library constructed from rat
                    bladder tissue. For a detailed description of the library
                    from which this clone was derived, please visit our web
                    site at ratest.eng.uiowa.edu. The subtraction has been
                    previously described in (Bonaldo, Lennon and Soares,
                    Genome Research 6:791-806, 1996)
                    TAG TISSUE=bladder
                    TAG_LIB=UI-R-DB0
                    TAG SEQ=AGC"
ORIGIN
                         25.0%; Score 31.2;
                                              DB 2;
                                                     Length 655;
 Query Match
 Best Local Similarity 53.2%; Pred. No. 28;
                                                 58;
           66; Conservative
                              0; Mismatches
                                                      Indels
                                                                    Gaps
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Qy
                       Db
         537 AAACAACAGTGACTTGTATCTAAGTCGTGTGGTGGAAGTGAAGACAGTGAAGACAATCCA 478
           62 CCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGCCAC 121
Qy
                                      - 11
                                            11
          477 ACAGCAGATAGAGGCTGGAAAGACCTTGTTATTTGATGTGATCTTAGGCAAAACAGCATG 418
Db
         122 TTTG 125
Qy
              +111
         417 TTTG 414
Db
RESULT 13
CW685817
                                    669 bp
                                              DNA
                                                               GSS 01-NOV-2004
LOCUS
            CW685817
                                                      linear
           OG_BBa0044A23.f OG_BBa Oryza glaberrima genomic clone OG_BBa0044A23
DEFINITION
            5', genomic survey sequence.
ACCESSION
            CW685817
VERSION
            CW685817.1 GI:55155379
KEYWORDS
           GSS.
SOURCE
            Oryza glaberrima (African rice)
 ORGANISM
           Oryza glaberrima
            Eukaryota; Viridiplantae; Streptophyta; Embryophyta; Tracheophyta;
            Spermatophyta; Magnoliophyta; Liliopsida; Poales; Poaceae;
            Ehrhartoideae; Oryzeae; Oryza.
REFERENCE
               (bases 1 to 669)
            Kim, H., Yu, Y., Wissotski, M., Byrne, M., Stum, D., Smart, D., Rao, K.,
 AUTHORS
            Luo, M., Jetty, R., Kudrna, D., Muller, C., Hatfield, J., Soderlund, C.
            and Wing, R.
  TITLE
            OMAP
  JOURNAL
            Unpublished (2004)
COMMENT
            Contact: Rod A. Wing
```

```
Arizona Genomics Institute
            University of Arizona
            Forbes Building Room 303, Tucson, AZ 85721-0036, USA
           Tel: 520 626 9595
            Fax: 520 621 1259
            Email: rwing@genome.arizona.edu
            PCR PRimers
            FORWARD: TAA TAC GAC TCA CTA TAG GG
           BACKWARD: CAC TCA TTA GGC ACC CCA
            Plate: 0044 row: A column: 23
           Seq primer: TAA TAC GAC TCA CTA TAG GG
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                     Location/Qualifiers
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ORIGIN
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                          59.8%; Pred. No. 33;
  Best Local Similarity
           52; Conservative
                                 0;
                                    Mismatches
                                                       Indels
                                                                     Gaps
                                                                             0;
            2 AAAAACAAAAAACCTTTACGTAACGTTTTTGCTGGGAGAAGACTACGAAGCACATTTT 61
Qy
              161 AGAAACAAACAAACTATAGGACATCTTTTTTTAGAGCGGGTAAAGTATGAAGGACTTTTG 220
Db
           62 CCAGGAAGTGTGGGCTGCAACGATTGT 88
Qy
                      \perp \perp \downarrow \downarrow
Db
          221 ACTGGGATCCTGTGGTAAAGCAATAGT 247
RESULT 14
BI527837/c
LOCUS
           BI527837
                                     422 bp
                                              mRNA
                                                       linear
                                                               EST 29-AUG-2001
           1024084H12.x1 C. reinhardtii CC-1690, normalized, Lambda Zap II
DEFINITION
           Chlamydomonas reinhardtii cDNA, mRNA sequence.
ACCESSION
           BI527837
VERSITON
           BI527837.1 GI:15368411
KEYWORDS
           EST.
SOURCE
           Chlamydomonas reinhardtii
  ORGANISM
           Chlamydomonas reinhardtii
           Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae;
           Chlamydomonadales; Chlamydomonadaceae; Chlamydomonas.
REFERENCE
           1 (bases 1 to 422)
 AUTHORS
           Grossman, A., Chang, C.-W., Davies, J., Harris, E., Hauser, C.,
           Lefebvre, P., McDermott, J.P., Shrager, J., Silflow, C. and Stern, D.
  TITLE
           Analyses of the Chlamydomonas reinhardtii Genome: A Model,
           Unicellular System for Analyzing Gene Function and Regulation in
            Vascular Plants. Project: 1024b
  JOURNAL
           Unpublished (2001)
COMMENT .
            Contact: Charles Hauser . .
            DCMB Box 91000
            Duke University
            Durham, NC 27708-1000
            Tel: 919 613 8159
```

Fax: 919 613 8177 Email: chauser@duke.edu. Location/Qualifiers **FEATURES** source 1. .422 /organism="Chlamydomonas reinhardtii" /mol type="mRNA" /strain="CC-1690 wild type mt+ 21gr" /db xref="taxon:3055" /clone_lib="C. reinhardtii CC-1690, normalized, Lambda Zap II" /note="Vector: pBluescript II SK-; Site 1: EcoRI; Site 2: XhoI; This library, constructed by John Davies and Jeffrey McDermott, combines cDNAs from CC-1690 cells grown to mid-log phase in TAP (acetate-containing) medium in the light, TAP medium in the dark, HS (minimal) medium in ambient levels of CO2 and HS medium bubbled with 5% CO2. PolyA mRNA was purified from each sample, pooled and cDNA synthesized. The cDNA was directionally cloned into lambda ZAP II (Stratagene) in the EcoRI (5') and XhoI (3') sites. pBluescript II SK- plasmids were excised from the lambda ZAP clones by superinfection with ExAssist (Stratagene) phage. The library was normalized using method 4 described in Bonaldo et al (1996) Genome Research 6: 791-806." ORIGIN . 24.6%; Score 30.8; DB 3; Length 422; Query Match Best Local Similarity 70.7%; Pred. No. 34; 0; Mismatches 0: 41; Conservative 17: Indels Gaps Qy 61 TCCAGGAAGTGTGGGCTGCAACGATTGTGCGCTCTTAACTAATCCTGAGTAAGGTGGC 118 264 TCATGGCAGCATGGGATGCACCGCGTGTGCGCTCTGCACACGTCGTGGGTGAGGTGGC 207 Db RESULT 15 BI999848/c BI999848 linear EST 25-OCT-2001 LOCUS 452 bp mRNA DEFINITION 1031079E03.x2 C. reinhardtii CC-1690, Stress II (normalized), Lambda Zap II Chlamydomonas reinhardtii cDNA, mRNA sequence. ACCESSION BI999848 BI999848.1 GI:16434622 VERSION KEYWORDS EST. SOURCE Chlamydomonas reinhardtii ORGANISM Chlamydomonas reinhardtii Eukaryota; Viridiplantae; Chlorophyta; Chlorophyceae; Chlamydomonadales; Chlamydomonadaceae; Chlamydomonas. REFERENCE (bases 1 to 452) **AUTHORS** Grossman, A., Chang, C.-W., Davies, J., Harris, E., Hauser, C., Lefebvre, P., McDermott, J.P., Shrager, J., Silflow, C. and Stern, D. Analyses of the Chlamydomonas reinhardtii Genome: A Model, TITLE Unicellular System for Analyzing Gene Function and Regulation in Vascular Plants. Project: 1031 JOURNAL Unpublished (2001) COMMENT Contact: Charles Hauser DCMB Box 91000 Duke University Durham, NC 27708-1000 Tel: 919 613 8159 Fax: 919 613 8177 Email: chauser@duke.edu.

Location/Qualifiers

FEATURES

```
1. .452
source
                /organism="Chlamydomonas reinhardtii"
                /mol type="mRNA"
                /strain="CC-1690 wild type mt+ 21gr"
                /db_xref="taxon:3055"
                /clone lib="C. reinhardtii CC-1690, Stress II
                (normalized), Lambda Zap II"
               /note="Vector: pBluescript II SK-; Site_1: EcoRI; Site_2:
               XhoI; Stress condition II library, constructed by John
               Davies and Jeffrey McDermott, combines cDNAs from CC-1690
               cells grown to mid-log phase in TAP (NH4+ - containing)
               and shifted to TAP - NO3- (24hrs); H2 production
               conditions (0, 12hr, 24hr) see Melis et al., (2000) Plant
               Phys. 122: 127-135; TAP + H2O2 (1, 12, 24 hr); TAP +
               sorbitol (1, 2, 6, 24 hr); TAP + Cd (1, 2, 6, 24 hr).
               PolyA mRNA was purified from each sample, pooled and cDNA
               synthesized. The cDNA was directionally cloned into lambda
               Zap II (Stratagene) in the EcoRI (5') and XhoRI (3')
               sites. pBluescript II SK- plasmids were excised from the
               lambda ZAP clones by superinfection with ExAssist
                (Stratagene) phage. The library was normalized using
               method 4 described in Bonaldo et al., (1996) Genome
               Research 6: 791-806."
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ORIGIN

24.6%; Query Match Score 30.8; DB 3; Length 452; 70.7%; Best Local Similarity Pred. No. 35; 41; Conservative 0; Mismatches Gaps 17; Indels 0;

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Search completed: April 25, 2006, 10:52:01

Job time: 1991 secs

SCORE 1.3 BuildDate: 12/06/2005